

Valneva SE

France / Biotechnology Euronext Paris Bloomberg: VLA FP ISIN: FR0004056851

Update

RATING PRICE TARGET

BUY € 8.10

Return Potential 181.6% Risk Rating High

STRONG Q1 RESULTS; WE SEE LYME MARKET PEAKING AT >USD2BN

Q1/25 product revenues of €48.6m (Q1/24: €32.4m) jumped 51.2% due largely to the absence of supply constraints which hampered sales of Ixiaro and third-party products in the prior year quarter. An improved gross margin and tight cost control lowered the operating cash outflow to €8.1m (Q1/24: €-28.4m). For the full year management is guiding towards an operating cash outflow of under €30m (FY/24: €67.2m). The end Q1/25 cash position was €153m, which in April was topped up by €14.2m in poceeds from an issue to the U.S. healthcare investor, Novo Holdings A/S. Management indicates that the cash runway extends until expected first commercial sales of the Lyme disease vaccine candidate, VLA15, in 2027. We think VLA15 will make Valneva profitable. We also expect commercialisation of VLA15 to trigger aggregate milestone payments of USD143m from partner Pfizer in 2027. Additionally, Valneva also stands to receive USD100m in commercial milestones based on cumulative VLA15 sales thresholds, and royalties of between 14% and 22%. No vaccine is currently available to prevent Lyme disease in humans. Pfizer and Valneva currently estimate the global market for a Lyme disease vaccine to exceed USD1bn. We see this figure as too conservative. With VLA15 phase 3 trial results expected by the end of this year, we have reworked our forecasts for the vaccine candidate. Based on eligible populations in endemic regions of the U.S. and EU of ca 70m and 162m respectively, an average price per shot of USD72, and penetration of 10%, we arrive at a peak value for the Lyme vaccine market of over USD2bn. An increase in our valuation of VLA15 outweighs a downward revision to our valuation of Valneva's chikungunya vaccine, IXCHIQ, prompted by slower than expected uptake. We raise our price target to €8.10 (previously: €7.70) and maintain our Buy recommendation. Upside: 182%.

(p.t.o.)

FINANCIAL HISTORY & PROJECTIONS

	2022	2023	2024	2025E	2026E	2027E
Revenue (€m)	361.3	153.7	169.6	186.1	203.3	466.7
Y-o-y growth	3.8%	-57.5%	10.3%	9.7%	9.3%	129.5%
EBIT (€m)	-113.4	-82.1	13.3	-59.9	-34.8	182.1
EBIT margin	n.a.	n.a.	7.9%	n.a.	n.a.	39.0%
Net income (€m)	-143.3	-101.4	-12.2	-79.0	-49.5	153.4
EPS (diluted) (€)	-1.24	-0.73	-0.08	-0.48	-0.30	0.92
DPS (€)	0.00	0.00	0.00	0.00	0.00	0.00
FCF (€m)	-274.7	-217.1	-83.7	-44.5	-21.8	151.1
Net gearing	-62.4%	64.5%	26.5%	76.9%	182.5%	-4.4%
Liquid assets (€m)	289.4	126.1	168.3	110.7	40.2	141.7

RISKS

Risks include, but are not limited to development, partnering, regulatory, competition and retention of key personnel.

COMPANY PROFILE

Valneva is a specialty vaccine company which develops and commercialises prophylactic vaccines for infectious diseases with significant unmet medical need. Valneva is currently commercialising three vaccines and has successfully advanced several vaccine candidates into and through the clinic, including candidates against Lyme disease, the chikungunya virus and COVID-19. Valneva is incorporated in France and had over 700 employees at end December 2024.

MARKET DATA	As of 20 May 2025
Closing Price	€ 2.88
Shares outstanding	167.27m
Market Capitalisation	€ 481.07m
52-week Range	€ 1.76 / 4.23
Avg. Volume (12 Months)	971,142

Multiples	2024	2025E	2026E
P/E	n.a.	n.a.	n.a.
EV/Sales	3.2	2.9	2.7
EV/EBIT	40.5	n.a.	n.a.
Div. Yield	0.0%	0.0%	0.0%

STOCK OVERVIEW



COMPANY DATA	As of 31 Mar 2025
Liquid Assets	€ 152.99m
Current Assets	€ 152.99m
Intangible Assets	€ 24.60m
Total Assets	€ 482.24m
Current Liabilities	€ 106.20m
Shareholders' Equity	€ 175.25m

SHAREHOLDERS

CDC	8.4%
Pfizer Inc.	5.7%
Novo Holdings A/S	4.8%
Groupe Grimaud la Corbiere	3.7%
Free Float and other	77.4%

Valneva SE

Figure 1: Q1/25 results versus consensus

€m	Q1/25	Q1/25- Consensus	Δ	Q1/24	Δ
Total revenues	49.2	41.0	20.0%	32.8	50.2%
Product revenues	48.6	39.1	24.3%	32.1	51.2%
Other revenues	0.6	1.6	-62.5%	0.6	-0.7%
R&D expenses	15.0	20.6	-27.2%	13.1	14.2%
SG&A	19.4	23.8	-18.5%	23.0	-15.7%
Adj. EBITDA	-0.6	-14.8	n.a.	73.0	n.a.

Source: Valneva

21 May 2025

Product sales 24% above consensus, but in line with management expectations As figure 1 above shows, product revenues jumped by 51.2%. Strong product revenue performance was helped by the absence of the supply constraints which crimped Q1/24 sales of Ixiaro and third party products. Q1/25 product sales were 24% above consensus and are equivalent to 28% of the midpoint of FY/25 product sales guidance of €170-€180m. However, CEO Thomas Lingelbach indicated during the analysts' call that the number was in line with management expectations on which FY/25 guidance is based.

Figure 2: Q1/25 product revenue breakdown

€m	Q1/25	Q1/24	Δ
lxiaro	27.5	16.6	65.7%
Dukoral	12.3	11.3	8.8%
IXCHIQ	3.0	0.2	1400.0%
Third party products	5.8	4.1	41.5%
Total product sales	48.6	32.1	51.2%

Source: Valneva

Q1/25 sales of Ixiaro were boosted by a strong increase in sales to the U.S. military following the signing of a new one-year USD32.8m contract with the U.S. Department of Defense at the end of January, and also helped by higher sales into travel channels and stock replenishment in indirect markets. The rise in Q1/25 Dukoral sales was driven by a shipment worth €1.1m following a cholera outbreak on the French island of Mayotte.

Ixiaro gross margin likely below Q1/25 level of 72.6% in coming quarters The gross margin on Ixiaro and Dukoral sales reached 72.6% (Q1/24: 51.8%) and 52.0% (Q1/24: 38.1%) respectively in Q1/25. The improvement was primarily due to better manufacturing performance with fewer batch failures and inventory adjustments. However, during the analysts' call CFO Peter Bühler stated that the Ixiaro gross margin is likely to be lower during the remaining quarters of 2025. The Dukoral gross margin is expected to be sustainable at the Q1/25 level.

IXCHIQ was first launched in the U.S. in Q1/24. Sales of the vaccine jumped to €3.0m in Q1/25 (Q1/24: €0.2m) as Valneva ramped up sales in the U.S. and also in France and Canada. The gross profit on Ixiaro sales in Q1/25 was €2.0m (equivalent to a margin of 67%). The Q1/24 gross profit was €-0.6m.

Take-up of IXCHIQ has been slower than Valneva's management initially expected In a presentation of July 2024 management held out the prospect of overall group product revenues doubling from the 2023 level of €145m by 2026. This guidance implied 2026 IXCHIQ sales of over €100m. However, in the 9M/24 report published last November, management announced a review of mid-term guidance for IXCHIQ after sales of the vaccine came in at a modest €1.8m for the first three quarters of 2024. To date Valneva has not published any updated mid-term guidance for IXCHIQ.

In our view initial sales of IXCHIQ have failed to meet expectations because of low awareness of chikungunya among travellers. A delay in publication in the U.S. of the ACIP's (Advisory Committee On Immunization Practices) recommendation on IXCHIQ in the influential Morbidity and Mortality Weekly Report has also slowed uptake.

FDA and EMA have suspended IXCHIQ recommendation for elderly Earlier this month the FDA and EMA (European Medicines Agency) both announced a temporary suspension of their recommendations for the use of IXCHIQ for individuals over 65 years old. The FDA and EMA took this step following reports of serious adverse events (SAEs) following vaccination with IXCHIQ in elderly people with significant underlying medical conditions and or co-medications. The EMA's decision was based on reports of 17 SAEs worldwide in elderly people (including two deaths). To date, over 40,000 doses of IXCHIQ have been used worldwide. Both the FDA and EMA have highlighted that the exact cause of these SAEs and their relationship with the vaccine has not yet been determined.

THE FDA/EMA may well rescind their recommendation suspensions over the next few weeks/months. However, the current FDA/EMA recommendation suspension will mean that 2025 IXCHIQ sales will be lower than they otherwise would have been, especially given that Valneva's marketing efforts had been particularly targeting elder adults, who are at higher risk for severe chikungunya infection than younger individuals.

Figure 3: IXCHIQ: Growing geographic footprint/progress in label extensions

Date	Event
November 2023	FDA approval for individuals 18 years of age and older
January 2024	First patient dosed in phase 2 pediatric trial In Dominican Republic and Honduras
May 2024	Further positive phase 3 data from adolescent phase 3 trial
June 2024	Canada - approval for individuals 18 years of age and older
July 2024	EU approval for individuals 18 years of age and older
September 2024	Adolescent label extension applications submitted to EU and Canada
November 2024	Adolescent label extension application submitted to FDA
December 2024	Exclusive License with Serum Institure of India
February 2025	UK approval for individuals 18 years of age and older
March 2025	Adolescent label extension application submitted to UK's MHRA
April 2025	EU approval for individuals 12 years of age and older
April 2025	Brazil - approval for individuals 18 years of age and older
May 2025	EU temporarily suspends use of IXCHIQ for individuals over 65 years of age
May 2025	U.S. CDC recommends pause in use of IXCHIQ for individuals over 65 years of age

Source: Valneva

We think South American/Asian chikungunya-endemic markets could generate USD250m of sales by 2030 Figure 3 illustrates IXCHIQ's growing geographic approval footprint and progress in label extensions since initial FDA approval in November 2023. Particularly noteworthy in our view are the April 2025 approval in Brazil – the first marketing authorisation for IXCHIQ in a chikungunya-endemic country – and the exclusive license with Serum Institute of India (SII) announced in December last year. SII is the world's largest manufacturer of vaccines by number of doses, and will work together with Valneva to bring IXCHIQ to the Indian market and certain other Asian countries, subject to local regulatory approvals. We believe the chikungunya-endemic markets in the low and medium income countries of South America and Asia could generate chikungunya vaccine sales of USD250m by 2030. Assuming a 50% market share for Valneva, and a 30% operating profit margin, implies an EBIT contribution of USD38m.

Q2/25 will benefit from large IXCHIQ order from La Réunion Valneva states that ca. 40,000 doses of IXCHIQ had been used by the end of Q1/25. So far this year the company has supplied 40,000 doses to La Réunion as the French island is currently undergoing a major chikungunya outbreak.

Only a small number of these 40,000 doses were supplied in Q1/25, suggesting that current quarter sales should beat the first quarter figure of €3.0m by a significant margin. We also expect the third and fourth quarters to be above the first quarter as recent geographic approvals and label extensions boost sales.

Pivotal results of the phase 3 trial of Valneva's Lyme disease vaccine candidate, VLA15, are expected by the end of this year. Based on what we know about the size of the populations in endemic areas of the U.S. and EU, and what we think are conservative pricing and penetration assumptions, we arrive at a peak value for the Lyme vaccine market of over USD2bn. This compares with Pfizer and Valneva's longstanding estimate of over USD1bn. We have summarised our assumptions in figure 4 below.

Figure 4: Lyme Disease vaccine market size assumptions

	U.S.	EU	Total
Endemic population (m)	87.0	202.0	289.0
Eligible population (m)	69.6	161.6	231.2
Total penetration	10%	10%	10%
No. individuals vaccinated (m)	7.0	16.2	23.1
Vaccination schedule (months)	primary:0,2,5-9,	primary: 0,2,5-9,	
vaccination schedule (months)	boosters:18 and then annually	boosters:18 and then annually	
Price per shot (USD)	100	60	72

Source: Valneva, First Berlin Equity Research estimates

The population living in areas of the U.S. and EU in which Lyme Disease is endemic are 87m and 202m respectively. 20% of these populations are ineligible for vaccination by virtue of being too young or too old. We estimate a penetration rate of 10%. If the authorities issue a blanket recommendation for individuals living in endemic areas, penetration could clearly be much higher. Our assumed vaccination schedule follows that used in the phase 3 trial plus annual boosters following the trial booster which is administered at 18 months. Valneva has not made any official statements on pricing. However, Pfizer CEO Albert Bourla has stated that no primary vaccination series (in this case 3 shots) should cost less than USD300. We have assumed USD100 per shot in the US and USD60 per shot in the EU. This implies an average price per shot of USD72.

Our long term assumption is that Valneva/Pfizer and Moderna each take 50% of the Lyme disease vaccine market Subject to positive results from the phase 3 trial, Pfizer expects to submit regulatory applications for VLA15 in the U.S. and Europe in 2026 and receive approval in 2027. Moderna is Valneva/Pfizer's main competitor in the development of a vaccine for Lyme Disease. Moderna expects to complete its parallel phase 1/2 studies of the Lyme disease vaccine candidates heptavalent mRNA-1975 and monovalent mRNA-1982 by 31 July this year. We would then expect a phase 3 trial over two tick seasons in 2026 and 2027, regulatory submission subject to a positive trial result in 2028, and approval in 2029. Over the medium term we assume that Valneva/Pfizer and Moderna each take 50% of the Lyme disease vaccine market.

We maintain our Buy recommendation and raise the price target from €7.70 to €8.10 Changes to our forecasts mainly reflect reductions in our IXCHIQ forecasts, and increases in our forecasts for VLA15 as explained above. We now see fair value for the Valneva share at €8.10 (previously: €7.70). We maintain our Buy recommendation. Upside: 182%.



Figure 5: Changes to our forecasts

€m		2025E			2026E			2027E	
EIII -	Old	New	Δ	Old	New	Δ	Old	New	Δ
Product revenues	203.7	176.1	-13.5%	290.1	200.3	-30.9%	355.4	333.7	-6.1%
of which:									
lxiaro	96.4	110.0	14.1%	101.9	121.0	18.8%	106.3	133.1	25.3%
Dukoral	34.7	36.3	4.8%	35.7	38.1	6.8%	36.8	40.0	8.9%
lxchiq	52.6	18.0	-65.8%	136.9	32.0	-76.6%	163.5	55.0	-66.4%
Lyme royalties/milestones	0.0	0.0	-	0.0	0.0	-	32.7	96.1	-
Third party revenues	20.1	11.8	-41.2%	15.7	9.2	-41.2%	16.1	9.5	-41.2%
Other revenue	10.3	10.0	-2.9%	10.6	3.0	-71.7%	140.9	133.0	-5.6%
Total revenues	214.0	186.1	-13.0%	300.7	203.3	-32.4%	496.3	466.7	-6.0%
Gross profit	115.0	92.6	-19.5%	174.5	121.0	-30.7%	353.2	355.3	0.6%
margin (%)	0.0	0.0	-	0.0	0.0	-	0.0	0.0	-
Sales & marketing	-57.0	-43.0	-	-72.5	-50.1	-	-88.9	-59.4	-
General & administrative	-53.0	-37.0	-	-55.1	-40.1	-	-60.4	-45.1	-
Research & development	-67.5	-95.0	-	-67.5	-80.1	-	-47.5	-83.2	-
Other income, net	20.0	22.5	12.5%	20.0	14.5	-27.5%	6.0	14.5	141.7%
EBIT	-42.5	-59.9	n.a.	-0.7	-34.8	n.a.	162.4	182.1	12.2%
margin (%)	-19.8%	-32.2%	-	-0.2%	-17.1%	-	32.7%	39.0%	-
Net financial result	-17.6	-17.6	-	-14.7	-14.7	-	-11.7	-11.7	-
EBT	-60.1	-77.5	n.a.	-15.4	-49.5	n.a.	150.7	170.5	13.1%
Tax	0.0	-1.5	-	0.0	0.0	-	-15.1	-17.0	-
Net income	-60.1	-79.0	n.a.	-15.4	-49.5	n.a.	135.7	153.4	13.1%
EPS (in EUR)	-0.43	-0.48	n.a.	-0.11	-0.30	n.a.	0.98	0.92	-6.1%
Adjusted EBITDA	-23.0	-39.1	n.a.	19.2	-13.5	n.a.	182.8	203.9	n.a.

Source: First Berlin Equity Research estimates

Figure 6: Valuation model

Compound	Project ¹⁾	Present Value	Market Size 2030	Market Share 2030	Sales 2030	PACME Margin ²⁾	Discount Factor	Time to Market
lxiaro	Japanese Encephalitis	€635.2M	€178.9M	90.0%	€1610M	40%	9.0%	-
Dukoral	Cholera & ETEC	€133.6M	€183.8M	25.0%	€45.9M	30%	90%	-
VLA15	Lyme Disease	€1,392.3M	€909.1M	261.6%	€2,378.6M	18%	10.0%	2 Years
VLA 1553	Chikungunya virus	€533.6M	€463.0M	50.0%	€231.5M	45%	9.0%	-
S4V	Shigellosis	€232.3M	n.a.	n.a	n.a	45%	10.0%	5 years
EB66 cell lir	ne Technology Platform	€11.9M			€19.7M	15%	9.0%	-
PACME PV		€2,938.9M						
Costs PV ³⁾		€1,732.6M						
NPV		€1,206.3M						
PV grants, o	collabs., 3rd party distrib.	€192.9M						
Proforma net cash		-€44.0M						
Fair Value		€1,355.2M						
Proforma share count (fully diluted)		167,272K						
Price Target		€8.10						

¹⁾ A project typically refers to a specific indication or, where necessary or relevant, a combination between indication and geographic market

Source: First Berlin Equity Research estimates

²⁾ PACME (Profit After Costs and Marketing Expenses) reflects the company's profit share on future revenues.

This share may be derived in the form of royalties (outsourced marketing/manufacturing) or operating EBITDA margin (in-house model), or some mix of both (depending on the specific parameters of partnership agreements)

³⁾ Includes company-level R&D, G&A, Financing Costs and CapEx; COGS and S&M are factored into the PACME margin for each project

Figure 7: Changes to our valuation model

	Old	New	Δ
PACME PV	€2,603.0M	€2,938.9M	12.9%
Costs PV	€1,523.1M	€1,732.6M	13.8%
NPV	€1,079.9M	€1,206.3M	11.7%
PV grants, collabs., 3rd party distrib., milestones	€192.1M	€192.9M	0.4%
Proforma net cash (inc. PRV)	-€23.2M	-€44.0M	89.5%
Fair Value	€1,248.8M	€1,355.2M	8.5%
Pro-forma share count	162,277K	167,272K	3.1%
Price Target	€7.70	€8.10	5.3%

Source: First Berlin Equity Research estimates

INCOME STATEMENT

All figures in EUR '000	2022	2023	2024	2025E	2026E	2027E
Product sales	114,797	144,624	163,253	176,100	200,319	333,714
Other income	246,506	9,089	6,326	10,000	3,000	133,000
Total revenues	361,303	153,713	169,579	186,100	203,319	466,714
Cost of materials/goods sold	-324,441	-100,875	-98,538	-93,545	-82,347	-111,376
Gross Profit	36,862	52,838	71,041	92,555	120,972	355,338
Sales & marketing	-23,509	-48,752	-52,356	-43,000	-50,080	-59,400
General & administrative	-34,073	-47,799	-42,750	-36,964	-40,064	-45,144
Research & development	-104,922	-59,894	-74,143	-95,000	-80,128	-83,160
Other operating items, net	12,199	21,520	111,538	22,500	14,500	14,500
Operating income (EBIT)	-113,443	-82,087	13,330	-59,909	-34,799	182,133
Net financial result	-18,794	-22,115	-21,623	-17,596	-14,714	-11,660
Foreign exchange gains/(loss)	-12,587	5,573	-3,193	0	0	0
Pre-tax income (EBT)	-144,815	-98,629	-11,486	-77,505	-49,513	170,473
Income taxes	1,536	-2,800	-761	-1,474	0	-17,047
Net income / loss	-143,279	-101,429	-12,247	-78,979	-49,513	153,426
EPS	-1.24	-0.73	-0.08	-0.48	-0.30	0.92
Adjusted EBITDA	-69,200	-65,187	32,930	-39,121	-13,532	203,893
Ratios as % of total revenues						
Gross margin	10.2%	34.4%	41.9%	49.7%	59.5%	76.1%
EBITDA margin	-19.2%	-42.4%	19.4%	-21.0%	-6.7%	43.7%
EBIT margin	-31.4%	-53.4%	7.9%	-32.2%	-17.1%	39.0%
Net margin	n.a.	n.a.	n.a.	n.a.	n.a.	32.9%
Expenses as % of total revenues						
Sales & marketing	-6.5%	-31.7%	-30.9%	-23.1%	-24.6%	-12.7%
General & administrative	-9.4%	-31.1%	-25.2%	-19.9%	-19.7%	-9.7%
Research & development	-29.0%	-39.0%	-43.7%	-51.0%	-39.4%	-17.8%
Y-Y Growth						
Product sales	82.3%	26.0%	12.9%	7.9%	13.8%	66.6%
Total revenues	3.8%	-57.5%	10.3%	9.7%	9.3%	129.5%
Operating income (ΕΒΠ)	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
Net income / loss	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.



All figures in EUR '000	2022	2023	2024	2025E	2026E	2027E
Assets						
Current Assets, Total	424,659	262,824	299,012	228,696	168,401	355,228
Cash and cash equivalents	289,430	126,080	168,271	110,709	40,196	141,651
Receivables	23,912	41,645	35,205	36,981	42,067	70,080
Inventories	35,104	44,466	53,662	40,503	40,064	66,743
Other current assets	76,213	50,633	41,874	40,503	46,073	76,754
Non-Current Assets, Total	196,685	197,238	201,020	195,307	188,763	186,098
Property, plant & equipment	112,435	136,198	138,883	137,874	136,834	135,763
Right of use assets	41,603	20,392	19,232	17,546	15,860	14,174
Intangibles	28,711	25,567	25,259	22,357	19,453	16,545
Equity-accounted investees	0	0	0	0	0	0
Other assets	8,299	8,489	8,041	7,925	7,011	10,011
Deferred tax assets	5,637	6,592	9,605	9,605	9,605	9,605
Total Assets	621,344	460,062	500,032	424,003	357,163	541,326
Shareholders' Equity & Debt						
Current Liabilities, Total	277,392	158,863	114,580	115,079	128,084	263,611
Short-term debt	11,580	44,079	20,852	31,189	35,195	110,463
Accounts payable	41,491	44,303	35,522	47,547	52,083	86,766
Other current liabilities and provisions	36,780	11,427	11,838	13,914	15,828	26,367
Current finance lease liabilities	25,411	2,879	2,508	2,288	2,068	1,848
Tax and employee-related liabilities	15,738	16,209	19,458	19,371	22,035	36,709
Current tax liability	532	632	1,742	770	875	1,458
Contract liabilities and refund liabilities	145,860	39,334	22,660	0	0	0
Longterm Liabilities, Total	124,155	172,952	204,199	188,656	158,202	53,295
Long term debt	87,227	132,768	166,521	145,658	110,463	0
Non-current finance lease liabilities	28,163	29,090	26,432	24,115	21,798	19,480
Other liabilities	1,436	1,153	593	8,355	15,535	23,525
Contract liabilities and refund liabilities	6,635	6,303	6,491	6,491	6,491	6,491
Shareholders Equity	219,797	128,247	181,253	120,269	70,877	224,420
Total Consolidated Equity and Debt	621,344	460,062	500,032	424,003	357,163	541,326
Dating						
Ratios Current ratio (x)	1.53	1.65	2.61	1.99	1.31	1.35
Quick ratio (x)	1.40	1.05	2.01	1.64	1.00	1.09
Net gearing	-62.4%	64.5%	26.5%	76.9%	182.5%	-4.4%
Book value per share (€)	1.31	04.5%	1.08	0.72	0.42	1.34
Net debt	-137,049	82,736	48,042	92,541	129,328	-9,859
Equity ratio	35.4%	27.9%	36.2%	28.4%	19.8%	41.5%
	JJ. 7/0	21.070	JU.2 /0	_0.470	10.070	11.070



CASH FLOW STATEMENT

All figures in EUR '000	2022	2023	2024	2025E	2026E	2027E
Net income / loss	-143,279	-101,429	-12,247	-78,979	-49,513	153,426
Depreciation and amortization	17,932	16,807	19,586	20,788	21,267	21,760
Impairment	0	0	0	0	0	0
Share-based payments	0	0	0	0	0	0
Gain from sale of priority review voucher	0	0	-90,833	0	0	0
Adjustments for non-cash transactions	44,070	44,984	48,979	46,858	42,981	57,467
Changes in non-current op. assets/lias.	-147,713	514	-178	0	0	0
Changes in w orking capital	1,732	-145,578	-11,394	4,014	95	-26,904
Other adjustments	0	0	0	0	0	0
Income tax	-154	-1,236	-1,545	-1,474	0	-17,047
Operating cash flow	-245,344	-202,745	-67,218	-29,582	-6,437	166,942
Property, plant and equipment	-29,246	-14,231	-13,865	-14,854	-15,299	-15,758
Investments in intangibles	-76	-81	-2,579	-100	-100	-100
Free cash flow	-274,666	-217,057	-83,662	-44,535	-21,836	151,084
Acquisitions & disposals, net	8	-7,482	165	0	0	0
Proceeds from sale of priority review vouche	0	0	90,833	0	0	0
Interest received	260	1,210	2,362	0	0	0
Investing cash flow	-29,054	-20,585	76,916	-14,954	-15,399	-15,858
Debt financing, net	37,538	79,014	-3,768	-13,063	-33,726	-37,732
Equity financing, net	189,837	-239	57,139	14,200	0	0
Payment of lease liabilities	-3,048	-3,127	-2,720	-237	-237	-237
Interest paid	-9,211	-12,567	-19,969	-17,596	-14,714	-11,660
Cash flow from financing	215,116	63,081	30,682	-16,696	-48,677	-49,629
Forex & other	2,026	-3,101	1,811	3,670	0	0
Net cash flows	-57,256	-163,350	42,191	-57,562	-70,513	101,455
Cash and equivs., start of the year	346,686	289,430	126,080	168,271	110,709	40,196
Cash and equivs., end of the year	289,430	126,080	168,271	110,709	40,196	141,651
Adj. EBITDA/share	-0.60	-0.47	0.22	-0.24	-0.08	1.22
Y-Y Growth						
Operating cashflow	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
Free cashflow a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
EBITDA/share a.	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.



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Category			2	
Current market	capitalisation (in €)	0 - 2 billion	> 2 billion	
Strong Buy ¹	An expected favourable price trend of:	> 50%	> 30%	
Buy	An expected favourable price trend of:	> 25%	> 15%	
Add	An expected favourable price trend of:	0% to 25%	0% to 15%	
Reduce	An expected negative price trend of:	0% to -15%	0% to -10%	
Sell	An expected negative price trend of:	< -15%	< -10%	

¹ The expected price trend is in combination with sizable confidence in the quality and forecast security of management.

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Report No.:	Date of publication	Previous day closing price	Recommendation	Price target
Initial Report	26 April 2017	€2.52	Buy	€4.00
238	↓	\downarrow	↓	↓
39	19 August 2022	€9.91	Add	€12.00
40	4 April 2023	€4.86	Buy	€8.90
41	22 May 2023	€5.69	Buy	€8.90
42	12 October 2023	€5.46	Buy	€8.90
43	27 February 2024	€3.21	Buy	€8.60
44	12 April 2024	€3.94	Buy	€8.60
45	20 August 2024	€3.78	Buy	€8.60
46	23 September 2024	€2.53	Buy	€7.70
47	Today	€2.88	Buy	€8.10

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