



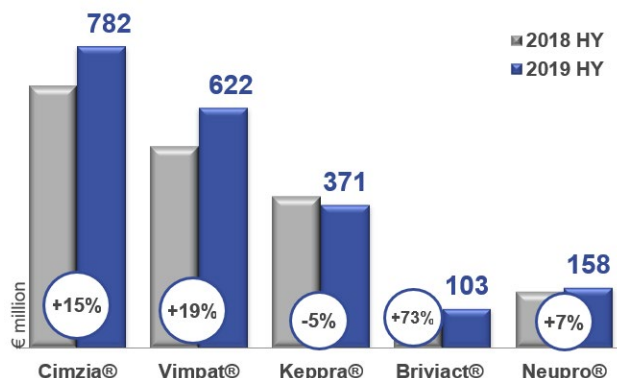
Brussels (Belgium), 25 July 2019 – 7:00 (CEST) – regulated information –
UCB Half Year Report 2019:

UCB's strong performance enables continued investment into future growth drivers

- Revenue reached € 2.3 billion (+2%, +4% CER¹)
 net sales increased to € 2.2 billion (+3%, +5% CER – adjusted² +11%, +7% CER)
- Underlying profitability (rEBITDA³) was € 724 million (-9%, -1% CER) or a ratio of 31%
- R&D update: Vimpat® with positive phase 3 results in PGTCS; four new phase 3 programs started: *bimekizumab* in psoriatic arthritis and axial spondyloarthritis; *padsevonil* in drug-resistant focal epilepsy; *rozanolixizumab* in myasthenia gravis
- Financial outlook for 2019 confirmed: Revenue expected to reach € 4.6 - 4.7 billion, recurring EBITDA³ should reach 27 - 29% of revenue, Core EPS of € 4.40 - 4.80 expected
- CFO transition in 2020

"The strong financial performance in the first six months 2019 enables us as planned to accelerate our investments into future growth drivers," said Jean-Christophe Tellier, CEO UCB. "Our ambition is to create patient value for specific populations. From our promising late-stage pipeline we had already two product approvals and started four new phase 3 programs. In this context, we are confirming our financial outlook for the full year 2019, while we also reconfirm our commitment to competitive profitability in the mid-term."

Core product net sales



Revenue for the first six months of 2019 reached € 2.3 billion (+2%; +4% at CER).

Net sales went up by 3% to € 2.2 billion (+5% CER), driven by the continued growth of UCB's core products.

Underlying profitability (rEBITDA³) reached € 724 million (-9%; -1% CER) due to higher operating expenses - reflecting the investments into the future of UCB, namely into product launches and product development.

Profit of the Group was to € 437 million of which € 411 million (-24%; -14% CER) is attributable to the UCB shareholders.

Core earnings per share⁴ were € 2.42 after € 3.09.

UCB's financial results HY 2019

€ million	2019 HY	2018 HY	Act	CER
Revenue	2 323	2 269	2%	4%
Net sales	2 219	2 146	3%	5%
rEBITDA ³	724	794	-9%	-1%
Number of shares (m)	187	188	-1%	
Core EPS ⁴ (€)	2.42	3.09	-22%	-12%

¹ CER = constant exchange rates

² Adjusted for divestitures and designated hedges reclassified to net sales

³ rEBITDA = recurring Earnings Before Interest, Taxes, Depreciation and Amortization charges

⁴ Core EPS = core earnings per share

R&D update

Neurology

In January 2019, **Vimpat® (lacosamide)** was approved in Japan for the treatment of partial onset seizures in children 4 years of age and older. In addition, two new formulations were approved: IV (intravenous) and dry syrup.

In June, the Vimpat® development program for the adjunctive treatment of primary generalized tonic-clonic seizures (PGTCS) in study participants 4 years of age and older achieved statistically significant positive results for both its primary (time to second seizure) and secondary efficacy (seizure freedom) endpoints. The novel primary endpoint “time-to-second-seizure” reduced placebo-exposure of patients substantially. Submissions of this new indication are planned in the first half 2020 to multiple regulatory agencies.

In March, UCB started an international (U.S., EU, Japan and China) phase 3 study with **padsevonil** in drug-resistant focal epilepsy patients. First headline results are expected in H2 2021 and will complement those from the ongoing Phase 2b, expected in H1 2020. Padsevonil is an innovative drug purposely designed with a novel dual mechanism of action to address the needs of uncontrolled patients.

In May, **Nayzilam® (midazolam)** nasal spray was approved in the U.S. to treat intermittent, stereotypic episodes of frequent seizure activity in people living with epilepsy. UCB expects to make the treatment available to patients in the coming months.

In March, UCB started as planned a phase 2, proof-of concept, study with its novel, subcutaneous FcRn (neonatal Fc receptor) monoclonal antibody, **rozanolixizumab**, in patients with chronic inflammatory demyelinating polyneuropathy (CIDP). First headline results are expected in H1 2021. In June, UCB started as scheduled the confirmatory study (phase 3) with **rozanolixizumab** in patients with myasthenia gravis (MG). First headline results are expected in H1 2021.

Immunology

In March, UCB announced the approval of **Cimzia® (certolizumab pegol)** in the U.S. to include a new indication for the treatment of adults with active non-radiographic axial spondyloarthritis (nr-axSpA) with objective signs of inflammation.

In July and in China, Cimzia® was approved in combination with methotrexate for the treatment of moderate to severe, active rheumatoid arthritis in adult patients.

In March and April, the second and the third phase 3 program with **bimekizumab** - in psoriatic arthritis and in axial Spondyloarthritis - were initiated, slightly earlier than planned. First topline results are expected at the end of 2021.

In June, UCB and its partner Biogen initiated preparations for a phase 3 program with **dapirolizumab pegol** in patients with active systemic lupus erythematosus (SLE) despite standard-of-care treatment. The program is expected to start in H1 2020. This decision is based on the promising results of the phase 2b clinical trial, of which interim results were presented at EULAR in June 2019.

Bone

Early January 2019, UCB and Amgen announced the approval of **Evenity® (romosozumab)** in Japan. Evenity® is approved to reduce the risk of fractures and increase bone mineral density in men and post-menopausal women with osteoporosis at high risk of fracture.

In April, Evenity® was approved in the U.S. for the treatment of osteoporosis in post-menopausal women at high risk for fracture.

In May, Evenity® was approved in South Korea and in June in Canada and Australia.

In June, the Committee for Medicinal Products for Human Use (CHMP) of the European Medicines Agency adopted a negative opinion for **romosozumab**. The companies seek the re-examination of the CHMP opinion.

Net sales break-down by core product⁵

€ million	2019 HY	2018 HY	Act	CER
U.S.	480	416	15%	8%
Europe	208	192	8%	8%
International markets	94	71	31%	32%
Total Cimzia®	782	679	15%	10%

Immunology

Cimzia® (certolizumab pegol) for patients living with autoimmune and inflammatory TNF mediated diseases, net sales went up to € 782 million, driven by continued, sustainable growth in all regions. Growth is also driven by new patient populations like women in child bearing age and people living with psoriasis.

€ million	2019 HY	2018 HY	Act	CER
U.S.	472	387	22%	14%
Europe	111	100	11%	11%
International markets	39	35	10%	6%
Total Vimpat®	622	522	19%	13%

Neurology: UCB's epilepsy franchise reached net sales of € 1.25 billion, a plus of 12%.

Vimpat® (lacosamide), with net sales of € 622 million, shows continued strong growth in all regions thanks to reaching more and more people living with epilepsy.

€ million	2019 HY	2018 HY	Act	CER
U.S.	103	99	4%	-3%
Europe	84	113	-26%	-26%
International markets	184	180	2%	1%
Total Keppra®	371	392	-5%	-8%

Keppra® (levetiracetam) for epilepsy, reported net sales of € 371 million, reflecting both, the strong, established brand and the maturity of the product. In Europe, Keppra® net sales were affected by a local, one-time rebate adjustment.

€ million	2019 HY	2018 HY	Act	CER
U.S.	81	46	76%	65%
Europe	19	13	55%	55%
International markets	3	1	> 100%	> 100%
Total Briviact®	103	60	73%	64%

Briviact® (brivaracetam), reached net sales of € 103 million. This is driven by significant growth in all regions where Briviact® is available to patients.

€ million	2019 HY	2018 HY	Act	CER
U.S.	46	41	13%	5%
Europe	83	85	-3%	-3%
International markets	29	22	36%	31%
Total Neupro®	158	148	7%	4%

Neupro® (rotigotine), the patch for Parkinson's disease, increased net sales to € 158 million, mainly in the U.S. and in international markets.

⁵ Due to rounding, some financial data may not add up in the tables.

HY 2019 financial highlights⁶

For the six months ended 30 June ¹ € million	Actual		Variance	
	2019	2018	Actual rates	CER
Revenue	2 323	2 269	2%	4%
Net sales	2 219	2 146	3%	5%
Royalty income and fees	33	56	- 41%	- 47%
Other revenue	71	67	6%	4%
Gross profit	1 725	1 696	2%	4%
Marketing and selling expenses	- 502	- 442	14%	10%
Research and development expenses	- 568	- 500	13%	12%
General and administrative expenses	- 96	- 88	8%	7%
Other operating income / expenses (-)	12	- 9	N/A	N/A
Recurring EBIT (rEBIT)	571	657	- 13%	- 3%
Non-recurring income/expenses (-)	27	19	47%	49%
EBIT (operating profit)	598	676	- 11%	- 1%
Net financial expenses (-)	- 53	- 46	17%	17%
Share of net profit of associates	- 1	- 1	4%	4%
Profit before income taxes	544	629	- 13%	- 3%
Income tax expense (-)	- 108	- 56	94%	95%
Profit from continuing operations	436	573	- 24%	- 14%
Profit/loss (-) from discontinued operations	1	1	0%	- 10%
Profit	437	574	- 24%	- 14%
Attributable to UCB shareholders	411	551	- 25%	- 15%
Attributable to non-controlling interests	26	23	15%	7%
Recurring EBITDA	724	794	- 9%	- 1%
Capital expenditure (including intangible assets)	194	265	- 27%	
Net financial debt ⁷	387	237	63%	
Operating cash flow from continuing operations	353	492	- 28%	
Weighted average number of shares - non-diluted (million)	187	188	- 1%	
EPS ⁸	2.20	2.93	- 25%	- 25%
Core EPS⁸	2.42	3.09	- 22%	- 12%

“The statutory auditor has issued an unqualified review report dated 24 July 2019 on the company’s condensed consolidated interim financial statements as of and for the six month period ended 30 June 2019, and has confirmed that the accounting data reported in the press release is consistent, in all material respects, with the accounts from which it has been derived.”

⁶ Due to rounding, some financial data may not add up in the tables.

⁷ For the net financial debt, the reporting date for comparative period is 31 December 2018.

⁸ € per weighted average number of shares - non diluted

Revenue and net sales in the first six months of 2019 reached € 2 323 million (+2%; +4% at CER) and € 2 219 million (+3%; +5% CER) respectively. Net sales before “designated hedging reclassified to net sales” (+ € 65 million in H1 2018 / - € 51 million in H1 2019) were up by 9% (+5% CER). Adjusted for divestitures in 2018, mainly “Innere Medizin”/Germany, and Q1 2019, the iron supplement Niferex® and before ‘designated hedges reclassified to net sales’ growth was +11%, underlying: +7% CER. This growth was driven by the continued positive performance of UCB’s core products.

Royalty income and fees decreased to € 33 million from € 56 million. Other revenue reached € 71 million after € 67 million.

Gross profit reached € 1 725 million reflecting a stable gross margin.

Operating expenses reached € 1 154 million (+11%; +8% CER) reflecting 14% higher marketing and selling expenses of € 502 million - driven by the launch of Cimzia® in psoriasis and launch preparations for Cimzia® in active non-radiographic axial spondyloarthritis (nr-axSpA), 13% higher research and development expenses of € 568 million – driven by the pipeline progress and resulting in a R&D ratio of 24% in the first six months of 2019 and 8% higher general and administrative expenses of € 96 million. This resulted in an operating expense ratio (in relation to revenue) of 50% after 46% in the first half 2018.

Underlying profitability – rEBITDA⁹ reached € 724 million after € 794 million (-9%; -1% CER) driven by continued net sales growth compensated by higher operating expenses ratio and reflecting the investments into the future of UCB, namely into product launches and product development. The recurring EBITDA ratio for the first six months of 2019 (in % of revenue) reached 31%, from 35% in 2018.

Non-recurring income was € 27 million after € 19 million income in 2018.

Net financial expenses increased to € 53 million from € 46 million.

Income tax expenses were € 108 million compared to € 56 million in June 2018. The average effective tax rate on recurring activities was 20% compared to 9% in the same period of last year, but in line with the 2018 full year effective tax rate. The low tax rate in the first six months 2018 was driven by phasing of expenses and the U.S. tax reform.

Profit of the Group amounted to € 437 million (after € 574 million), of which € 411 million is attributable to UCB shareholders and € 26 million (after € 23 million) to non-controlling interests.

Core earnings per share, which reflect profit attributable to UCB shareholders, adjusted for the after-tax impact of non-recurring items, the financial one-offs, the after-tax contribution from discontinued operations and the net amortization of intangibles linked to sales, reached € 2.42 after € 3.09 based on 187 million weighted average shares outstanding.

Outlook 2019 confirmed - UCB confirms its expectations for 2019 revenue to reach approximately € 4.6 – 4.7 billion, recurring EBITDA in the range of 27 - 29% of revenue and core earnings per share are therefore expected in the range of € 4.40 – 4.80 based on an average of 188 million shares outstanding.

⁹ rEBITDA = recurring Earnings Before Interest, Taxes, Depreciation and Amortization charges

CFO transition in 2020 - Five consecutive years of growth and strong half-year results in 2019 have underscored UCB's robust performance over time. UCB is poised to further accelerate and expand to deliver on its Patient Value ambition, potentially launching 6 new products in the next 5 years.

With this goal in mind, UCB continues to focus its energies on strengthening its operating model to ensure maximum agility to meet our growth expectations for the years ahead. As part of this evolution, UCB implements in the coming months changes to its Executive Committee's structure. These changes will involve adjusting the size of the Committee; allowing it to enhance the focus on the company's core activity areas with increased agility. This will enable UCB to further evolve its patient value strategy.

In this context, UCB also announces that, after close to 13 years in the role, the company's Chief Financial Officer, Detlef Thielgen, will be transitioning out within a year from now. A search for a successor has just begun.

CEO Jean-Christophe Tellier said: "During his 13 years at UCB, Detlef has played an instrumental part in UCB's success and in bringing the company to the strong position it enjoys today. On behalf of the Board, of the Executive Committee and of all UCB teams, I would like to thank Detlef for his dedication to UCB over the years and for successfully helping to lead the company through its different transformation phases. Detlef and I will continue to work closely together to drive UCB forward, deliver on our objectives and ambition and ensure a smooth transfer of responsibilities and knowledge to his future successor."

Find the HY financial reports on UCB website: <http://www.ucb.com/investors/Download-center>

Today, UCB will host a conference call/video webcast at 08.00 (EDT) / 13.00 (BST) 14.00 (CEST).
Details are available on <https://www.ucb.com/investors/UCB-financials/Half-year-financial-results>.

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About UCB

UCB, Brussels, Belgium (www.ucb.com) is a global biopharmaceutical company focused on the discovery and development of innovative medicines and solutions to transform the lives of people living with severe diseases of the immune system or of the central nervous system. With more than 7 500 people in approximately 40 countries, the company generated revenue of € 4.6 billion in 2018. UCB is listed on Euronext Brussels (symbol: UCB). Follow us on Twitter: @UCB_news

Forward looking statements

This press release contains forward-looking statements based on current plans, estimates and beliefs of management. All statements, other than statements of historical fact, are statements that could be deemed forward-looking statements, including estimates of revenues, operating margins, capital expenditures, cash, other financial information, expected legal, political, regulatory or clinical results and other such estimates and results. By their nature, such forward-looking statements are not guarantees of future performance and are subject to risks, uncertainties and assumptions which could cause actual results to differ materially from those that may be implied by such forward-looking statements contained in this press release. Important factors that could result in such differences include: changes in general economic, business and competitive conditions, the inability to obtain necessary regulatory approvals or to obtain them on acceptable terms, costs associated with research and development, changes in the prospects for products in the pipeline or under development by UCB, effects of future judicial decisions or governmental investigations, product liability claims, challenges to patent protection for products or product candidates, changes in laws or regulations, exchange rate fluctuations, changes or uncertainties in tax laws or the administration of such laws and hiring and retention of its employees.

Additionally, information contained in this document shall not constitute an offer to sell or the solicitation of an offer to buy any securities, nor shall there be any offer, solicitation or sale of securities in any jurisdiction in which such offer, solicitation or sale would be unlawful prior to the registration or qualification under the securities laws of such jurisdiction. UCB is providing this information as of the date of this document and expressly disclaims any duty to update any information contained in this press release, either to confirm the actual results or to report a change in its expectations.

There is no guarantee that new product candidates in the pipeline will progress to product approval or that new indications for existing products will be developed and approved. Products or potential products which are the subject of partnerships, joint ventures or licensing collaborations may be subject to differences between the partners. Also, UCB or others could discover safety, side effects or manufacturing problems with its products after they are marketed.

Moreover, sales may be impacted by international and domestic trends toward managed care and health care cost containment and the reimbursement policies imposed by third-party payers as well as legislation affecting biopharmaceutical pricing and reimbursement.