

Pharming Group NV

Half Year Results 2017

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Safe harbour statement

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Half year results 2017

Operational highlights

- Accelerated the delivery of operating profitability as a result of strategic decision to reacquire commercial rights to RUCONEST® in North America
- The integration of the US RUCONEST® business into Pharming is firmly on track, with full transition of sales and marketing from Valeant Pharmaceuticals International completed
- Significant Investment made in a full US commercialization team
- Positive EMA amendment to the marketing authorization in Europe to allow self-administration of RUCONEST® for HAE attacks with a new custom-designed RUCONEST® Administration Kit
- Successfully refinanced company debt on more favourable commercial terms in order to recover dilutive share capital and release short and medium term cash to invest in accelerating commercialization efforts

Financial highlights

- Revenues from product sales for the half year increased by 617% to €30.1 million (HY 2016: €4.2 million), as a result of the combined effect of receiving all of the revenue from US product sales (instead of the previous 30% supply share of net sales), increasing patient numbers and product sales
- Total revenues increased by 477% to €30.6 million (including €0.5 million of license revenue) from €5.3 million (including €1.1 million in license revenue) in HY 2016
- Operating results improved to a profit of €4.2 million from a loss of €6.2 million in HY 2016, despite a considerable increase in commercialization activities, especially in the US
- The net result was a loss of €30.2 million (HY 2016: loss of €6.7 million), mainly as a result of one-off financing expenses required to be shown under IFRS associated with the extinction of the Amortising Convertible Bonds 2017/2018 and replacement of the previous debt facility
- The company's cash position decreased from €32.1 million at year-end 2016 to €25.2 million at 30 June 2017 (up from €21.7 million at 30 June 2016), largely due to pending trade payments under the Valeant transition services agreement

Long-term growth strategy

Investment in infrastructure to drive long-term growth

In the US

- Acquired entire Valeant sales team as part of transaction
- Expanded sales team and management, led by former senior HAE commercial executive as VP Commercial Operations
- Medical Science Liaison, Patient Services, Market Access and Managed Care teams in place from start of May
- Major overhaul of Positioning, Messages, Business Rules and Operating policies, and re-launch of full service patient care program RUCONEST[®] SOLUTIONS
- Experienced Commercial Advisory Board determining and monitoring strategy in the US, chaired by former CEO of a NASDAQ 100 Biotech and including former leading senior HAE commercial executives

In the EU

- Administration Kit now available for use in various EU markets, following approval of the Educational Materials by the local authorities in those markets

Next generation RUCONEST® - improving convenience

- RUCONEST® efficacy and safety profile for treatment of acute attacks is unsurpassed (on the basis of comparing published literature and patient experience)
- Next step: improving convenience of use
 - New highly concentrated vial in development for faster application of IV therapy (significantly lower volume and very rapid dissolution)
 - New vial will also enable clinical trials to test sub-cutaneous (SC) and intra-muscular (IM) injections for both acute and prophylaxis of HAE attacks
- Clinical trials for SC and IM applications are planned to start in early 2018

Financial Statements

Refinancing on improved commercial terms

- Recently completed \$100m 4 year debt facility (12 + 36) with OrbiMed
 - Interest approximately 12%, reducing to 11% if the company reaches \$100m in sales
 - Replaces original \$40 million of debt and remainder of 18 months €45 million amortising convertible bonds
 - Cash burn to be reduced by €16m in 2017, and €8m in 2018, due to lower repayments on debt and amortising bonds and lower cash interest and
 - Recovery of 115 million shares (24% of outstanding shares) which would otherwise have been issued at prices far below the current share price
- €11.8 million (\$13 million) unsecured tradeable 5 year (2016-2021) 8.5% convertible bonds with a conversion price of €0.284 now listed on Cayman exchange CSX

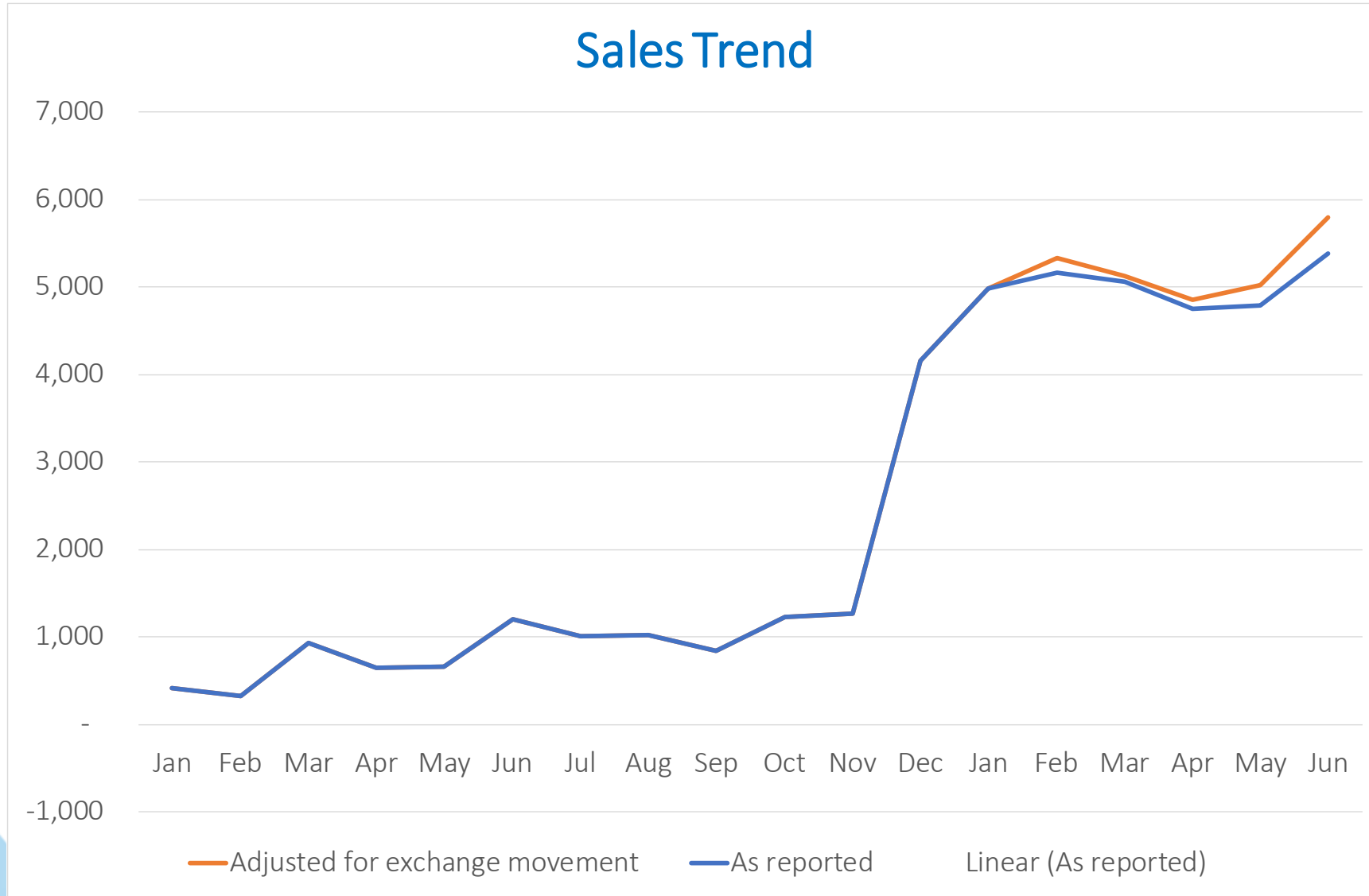
Income statement

Amounts in €'000	HY 2017	HY 2016
Product sales	30,109	4,170
Release of deferred license fee income	536	1,104
Revenues	30,645	5,274
Costs of product sales	(3,745)	(1,795)
Inventory impairments	88	(209)
Costs of sales	(3,657)	(2,004)
Gross profit	26,988	3,270
Other income	167	195
Research and development	(9,154)	(7,029)
General and administrative	(2,628)	(2,049)
Marketing and sales	(11,140)	(598)
Costs	(22,922)	(9,676)
Operating result	4,233	(6,211)

Monthly net revenues from sales 2016-H1 2017

Sales Trend

€'000



Adjusting for the effect of exchange rate movements, underlying Q2 sales were higher than Q1 (€15.7m v € 15.4m)

Income statement

Amounts in €'000, except per share data	HY 2017	HY 2016
Operating result	4,233	(6,211)
Fair value gain/(loss) on revaluation derivatives	(1,225)	455
Other financial income and expenses	(33,226)	(978)
Financial income and expenses	(34,451)	(523)
Result before income tax	(30,218)	(6,734)
Income tax expense	-	-
Net result for the period	(30,218)	(6,734)
Attributable to:		
Owners of the parent	(30,218)	(6,734)
Total net result	(30,218)	(6,734)
Basic earnings per share (€)	(0.063)	(0.016)

Balance sheet - Assets

Amounts in €'000	30 June 2017	31 December 2016
Intangible assets	55,855	56,680
Property, plant and equipment	7,104	6,043
Long term prepayment	2,644	1,622
Restricted cash	248	248
Non-current assets	65,851	64,593
Inventories	17,473	17,941
Trade and other receivables	18,645	12,630
Cash and cash equivalents	24,997	31,889
Current assets	61,115	62,190
Total assets	126,966	126.783

Balance Sheet - Liabilities

Amounts in €'000	30 June 2017	31 December 2016
Share capital	4,839	4,556
Share premium	310,907	301,876
Legal reserves	(612)	60
Accumulated deficit	(308,370)	(279,025)
Shareholders' equity	6,764	27,467
Loans and borrowings (more than one year)	78,628	40,395
Deferred license fees income	1,867	2,270
Finance lease liabilities	572	599
Other provisions	4,674	4,674
Non-current liabilities	85,741	47,938
Loans and borrowings (less than one year)	11,028	26,136
Deferred license fees income	811	943
Derivative financial liabilities	7,354	9,982
Trade and other payables	15,002	14,054
Finance lease liabilities	266	263
Current liabilities	34,461	51,378
Total equity and liabilities	126,966	126.783

Cash flow

Amounts in €'000	HY 2017	HY 2016
Operating result	4,233	(6,211)
Non-cash adjustments:		
Depreciation, amortization	1,689	316
Accrued employee benefits	872	914
Deferred license fees	(536)	(1,104)
Operating cash flows before changes in working capital	6,258	(6,084)
Changes in working capital:		
Inventories	468	(3,132)
Trade and other receivables	(6,015)	(2,330)
Payables and other current liabilities	(1,792)	3,214
Total changes in working capital	(7,339)	(2,247)
Changes in non-current assets, liabilities and equity	(3,109)	(258)
Net cash flows used in operating activities	(4,190)	(8,590)

Cash flow

Amounts in €'000	HY 2017	HY 2016
Net cash flows used in operating activities	(4,190)	(8,590)
Capital expenditure for property, plant and equipment	(1,457)	(752)
Investment intangible assets	(598)	-
Net cash flows used in investing activities	(2,055)	(752)
Proceeds of debt loans and borrowings	89,139	-
Payments of transaction fees	(16,051)	-
Repayments and interest on loans	(73,399)	(536)
Proceeds of equity and warrants	284	-
Net cash flows from financing activities	(27)	(536)
Increase (decrease) of cash	(6,272)	(9,878)
Exchange rate effects	(620)	(293)
Cash and cash equivalents at 1 January	32,137	31,843
Total cash at 30 June	25,245	21,672
Of which restricted cash	248	270
Cash and cash equivalents at 30 June	24,997	21,402

Outlook 2017

Outlook for remainder 2017

- Increasing sales and continued positive operating results
- Investment in the production of RUCONEST® in order to ensure continuity of supply.
- Assessment of the clinical trial results for RUCONEST® in prophylaxis of HAE by the US FDA and the development of other versions of RUCONEST®
- Increasing marketing activity where this can be profitable for Pharming, in addition to our current territories of Austria, France, Germany, United Kingdom and the Netherlands
- Continue to support our marketing partners in order to maximize the sales and distribution potential of RUCONEST® for patients in all territories, as we continue to believe that RUCONEST® represents a fast, effective, reliable and safe therapy option for HAE patients
- Continue to invest in the new pipeline programs in Pompe Disease and Fabry Disease

Tickers:
ENXTAM: PHARM
Bloomberg: PHAR.AS