



Pharming Group NV

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Interim Financial Results***

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Operational Highlights

- Rollout of Ruconest® in Europe progressing well
 - Q2, 2011 sales up 26% to €0.2 million
 - Reimbursement at national & local levels ongoing
- New data announced supporting efficacy of Rhucin® across all HAE attack locations
 - (7th C1INH deficiency workshop, Budapest, Hungary)
- Commercialization agreement signed with Megapharm for Ruconest® in Israel
- Protocol for Phase III study 1310 of Rhucin® amended
 - Primary endpoint modified
 - Increased patient numbers
 - Completion expected by Q3, 2012
- SPA agreement reached with the FDA on Phase III study 1310, announced earlier today
- Granting of US patent further extends protection of core technology platform

H1 2011 Overview

- €10.0 million received from Socius
 - €9.0 million received
 - plus €1.0 million from exercise of warrants
- DNage entered liquidation & subsequently deconsolidated
 - Curtails cash burn
 - €1.6 million operating cash outflow in HY1 2010 (nil in 2011)
- Entered financial lease for portion of new assets in use by Sanofi
 - Type of non dilutive funding

Financial Highlights

- Revenues & other income increased to €1.4 million (H1 2010: €0.1m)
- Operating costs decreased to €9.0 million (H1 2010: €10.1m)
 - Cash outflows of €8.9m (H1 2010: €8.4m ex. DNage)
 - but 2010 period included significant partnering cash income
- Significant reduction in net loss to €8.0 million (H1 2010: €28.0m)
 - Includes one-time €0.6m profit on discontinued operations (H1 2010: €1.7m loss)
- Cash at June 30, 2011 of €11.0 million (FY 2010: €10.5m)
 - Strengthened post period by private placement (€3.2m)

Deliverables

- EU & US commercialisation deals signed 
- EU launch of Ruconest® 
- Progress on US development pathway for Rhucin® 
- Continue EU rollout of Ruconest® **ongoing**
- Expand geographic coverage of Rhucin® franchise  **ongoing**
- Expand rhC1-INH platform 
 - Initiate reperfusion injury programmes **ongoing**
- Leverage potential of the platform **ongoing**
 - New proteins, new indications

Upcoming Milestones

- Continue EU rollout of Ruconest ®
- Expand geographic coverage of Rhucin® franchise
 - ROW Business Development initiatives on going
- Expand rhC1-INH platform
 - AMR ongoing
 - Ischaemia Reperfusion Injury evaluation ongoing
- Leverage potential of the platform
 - Follow up ongoing

Summary

Significant progress made during the period

Commercialisation

- Rollout of Ruconest® in Europe is progressing well
- Reimbursement at national & local levels is ongoing
- Commercialization agreement with Megapharm for Ruconest® in Israel
 - represents first step in global expansion of C1 Inhibitor franchise

Regulatory

- Clarity from the FDA on US regulatory pathway for Rhucin®
- Agreement with FDA on SPA reached

Financials

- Significant revenue growth and net loss reduction
- Increased cash position
- €10.0 million received from Socius

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