

23 September 2010

'RESEARCH NEVER STOPS'

info@evotec.com | www.evotec.com

For further information,
please contact

Dr Werner Lanthaler
Chief Executive Officer
+49.(0)40.560 81-242
+49.(0)40.560 81-333 Fax
werner.lanthaler@evotec.com

Evotec AG
Schnackenburgallee 114
22525 Hamburg (Germany)

Evotec Signs Ion Channel Hit Identification Agreement with Almirall

Hamburg, Germany – 23 September 2010: Evotec AG (Frankfurt Stock Exchange: EVT, TecDAX) today announced that it has entered into a collaboration with Almirall S.A. (BMAD: ALM), to identify small molecule modulators of an ion channel target, selected by Almirall, involved in respiratory diseases.

Evotec will apply its in-depth electrophysiology and ion channel pharmacology expertise, as well as its state-of-the-art screening platform, for the identification and validation of novel modulators of the selected ion channel.

Dr Jorge Beleta, Director Discovery Strategy and Alliances of Almirall, commented: "We are impressed not only by Evotec's ion channel drug discovery platform but more importantly by their track record of success and scientific understanding of this challenging class of targets. We believe that the Evotec technology and scientists will provide Almirall with the best chances of finding novel, high quality hits, ready for optimisation."

Dr Mario Polywka, COO of Evotec stated: "We are proud to have been selected by Almirall for this important project. We have invested significantly in our ion channel and electrophysiology platform and scientists in recent years, and this programme is further validation of the value we can bring to our partners. We look forward to collaborating with and supporting Almirall in their quest to find novel treatments to address respiratory diseases."

Evotec has a unique assay development and screening platform, built around proprietary and the latest commercial technologies, providing a flexible and high quality approach to lead identification for ion channels and other target classes.

No financial details are disclosed.

ABOUT EVOTEC AG

Evotec is a leader in the discovery and development of novel small molecule drugs with operational sites in Europe and Asia. The Company has built substantial drug discovery expertise and an industrialised platform that can drive new innovative small molecule compounds into the clinic. In addition, Evotec has built a deep internal knowledge base in the treatment of diseases related to neuroscience, pain, oncology, inflammation and metabolic diseases. Leveraging these skills and expertise the Company intends to develop best-in-class differentiated therapeutics and deliver superior science-driven discovery alliances with pharmaceutical and biotechnology companies. Evotec has long-term discovery alliances with partners including Boehringer Ingelheim, CHDI, Genentech, Novartis, Ono Pharmaceutical and Roche. Evotec has product

candidates in clinical development and a series of preclinical compounds and development partnerships, including for example a strategic alliance with Roche for the EVT 100 compound family, subtype selective NMDA receptor antagonists for use in treatment-resistant depression and an alliance in the field of diabetes with Andromeda (Teva). For additional information please go to www.evotec.com.

ABOUT ALMIRALL

Almirall is an international pharmaceutical company based on innovation and committed to health. Headquartered in Barcelona, Spain, it researches, develops, manufactures and commercialises its own R&D and licensed drugs with the aim of improving people's health and wellbeing.

Almirall focuses its research resources on therapeutic areas related to the treatment of asthma, COPD (Chronic Obstructive Pulmonary Disease), rheumatoid arthritis, multiple sclerosis, psoriasis and other dermatological conditions.

Almirall's products are currently present in over 70 countries while it has direct presence in Europe and Latin America through 12 affiliates.

For further information please visit the website at: www.almirall.com

FORWARD-LOOKING STATEMENTS — *Information set forth in this press release contains forward-looking statements, which involve a number of risks and uncertainties. Such forward-looking statements include, but are not limited to, statements about our 2010 financial outlook and our expected financial results in future quarters, our revised revenue guidance for 2010 and expected revenue growth, our ability to deliver on our liquidity guidance, our belief that we are on course to sustainable profitability latest in 2012, our expectations and assumptions concerning regulatory, clinical and business strategies, the progress of our clinical development programmes and timing of the commencement and results of our clinical trials, strategic collaborations and management's plans, objectives and strategies. These statements are neither promises nor guarantees, but are subject to a variety of risks and uncertainties, many of which are beyond our control, and which could cause actual results to differ materially from those contemplated in these forward-looking statements. In particular, the risks and uncertainties include, among other things: risks that product candidates may fail in the clinic or may not be successfully marketed or manufactured; the risk that we will not achieve the anticipated benefits of our collaborations, partnerships and acquisitions in the timeframes expected, or at all; risks relating to our ability to advance the development of product candidates currently in the pipeline or in clinical trials; our inability to further identify, develop and achieve commercial success for new products and technologies; the risk that competing products may be more successful; our inability to interest potential partners in our technologies and products; our inability to achieve commercial success for our products and technologies; our inability to protect our intellectual property and the cost of enforcing or defending our intellectual property rights; our failure to comply with regulations relating to our products and product candidates, including FDA requirements; the risk that the FDA may interpret the results of our studies differently than we have; the risk that clinical trials may not result in marketable products; the risk that we may be unable to successfully secure regulatory approval of and market our drug candidates; and risks of new, changing and competitive technologies and regulations in the U.S. and internationally.*

The list of risks above is not exhaustive. Our most recent Annual Report on Form 20-F, filed with the Securities and Exchange Commission, and other documents filed with, or furnished to the Securities and Exchange Commission, contain additional factors that could impact our businesses and financial performance. We expressly disclaim any obligation or undertaking to release publicly any updates or revisions to any such statements to reflect any change in our expectations or any change in events, conditions or circumstances on which any such statement is based.