

Almirall and Forest to present data from phase III studies of acclidinium bromide in Chronic Obstructive Pulmonary Disease (COPD)

- **Data to be presented at the 2011 American Thoracic Society (ATS) International Conference**

Barcelona, May 9th, 2011-- Almirall, S.A. (ALM.MC) and Forest Laboratories, Inc. (NYSE: FRX) today announced that they are presenting additional data from ATTAIN, a pivotal Phase III study comparing the efficacy and safety of inhaled acclidinium bromide 200µg and 400µg twice daily (BID) versus placebo in patients with moderate to severe COPD, at the annual ATS 2011, the annual international conference of the American Thoracic Society taking place in Denver May 13-18, 2011. Positive top-line results from this six month double-blind placebo-controlled study were first reported in January 2011.

Additional detail from the ATTAIN study, which was accepted as a late-breaker, will be presented in poster format on Sunday May 15th (8:15 am - 4:30 pm, Session A45, D100, by PW Jones).

Three other acclidinium clinical posters will also be presented at the ATS on Sunday May 15th. These posters include additional data from the Phase III ACCORD COPD I study regarding the effects of acclidinium therapy on symptoms and quality of life, as well as rescue medication use (8:15 am - 4:30 pm, Session A45, D75; EM Kerwin, D97 by A D'Urzo, D99 by AF Gelb).

In addition, results from one pharmacokinetic study with acclidinium BID (Sunday, May 15 at 8:15 am - 4:30 pm, Session A45, D98; K Lasseter), and five preclinical studies (Sunday, May 15 Session A45, D66; J Milara and Monday, May 16 Session B69, L21; J Milara, L30; J Milara, L29; D Dominguez-Fandos, L23; E Ferrer) will also be presented.

About ATTAIN Phase III study

ATTAIN (**A**clidinium **T**o **T**reat **A**irway obstruction **I**n **C**OPD **p**atie**N**ts) was conducted in Europe and South Africa. It was a 24 week study, which assessed the long term bronchodilator efficacy and safety of inhaled acclidinium bromide 200 µg and 400 µg, both administered BID, compared to placebo in 828 moderate to severe COPD patients (mean baseline FEV1= 1480mL). In addition, it assessed the benefits of acclidinium bromide 200µg and 400µg, compared to placebo, in disease-related health status and COPD symptoms.

About acclidinium bromide and the Genuair[®] inhaler

Acclidinium bromide is a novel, long-acting inhaled anticholinergic bronchodilator which has a long residence time at the M3 receptors and a shorter residence time at the M2 receptors. Acclidinium is rapidly hydrolyzed in human plasma to two major inactive metabolites. Almirall licensed Forest Laboratories, Inc. the US rights for acclidinium as well as to Kyorin exclusive rights to develop and commercialize acclidinium bromide in Japan, while Almirall maintains rights for the rest of the world. The companies are jointly involved in the development of the compound.

Acclidinium bromide is administered to patients using a novel, investigational, state-of-the-art multidose dry powder inhaler (MDPI), Genuair[®]. The Genuair[®] inhaler was designed with a feedback system, which through a 'colored control window' and an audible click helps confirm that the patient has inhaled correctly. It contains multiple doses of acclidinium, includes a visible dose-level indicator and also incorporates safety features such as an anti-double-dosing mechanism

and an end-of-dose lock-out system to prevent use of an empty inhaler. Genuair[®] is a registered trademark owned by Almirall, S.A.

About COPD

The World Health Organization (WHO) has described COPD as a global epidemic; an estimated 210 million people have COPD worldwide and more than 3 million people died of the condition in 2005, which is equal to 5% of all deaths globally that year. Total deaths from COPD are projected to increase by more than 30% in the next 10 years without interventions to cut risks, particularly exposure to tobacco smoke.

In patients with COPD the airways in the lungs typically lose their elasticity, produce excess mucus and become thick and inflamed, limiting the passage of air. The most common symptoms of COPD are breathlessness (or a "need for air"), abnormal sputum (a mix of saliva and mucus in the airway), and a chronic cough. Daily activities, such as walking up a short flight of stairs or carrying a suitcase, can become very difficult as the condition gradually worsens. There are significant unmet needs in the treatment of COPD including limited therapeutic options to improve lung function, reduce symptoms and control exacerbations.

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), gastrointestinal disorders, 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: www.almirall.com.

About Forest Laboratories

Forest Laboratories' (NYSE: FRX) longstanding global partnerships and track record developing and marketing pharmaceutical products in the United States have yielded its well-established central nervous system and cardiovascular franchises and innovations in anti-infective medicine. The Company's pipeline, the most robust in its history, includes product candidates in all stages of development across a wide range of therapeutic areas. The Company is headquartered in New York, NY. To learn more, visit www.FRX.com.

Except for the historical information contained herein, this release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. These statements involve a number of risks and uncertainties, including the difficulty of predicting FDA approvals, the acceptance and demand for new pharmaceutical products, the impact of competitive products and pricing, the timely development and launch of new products, and the risk factors listed from time to time in Forest Laboratories' Annual Report on Form 10-K, Quarterly Reports on Form 10-Q, and any subsequent SEC filings.

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