



BIOALLIANCE PHARMA TO COMPLETE NDA FOR LORAMYC[®] WITH DATA ON DEBOSSSED MUCOADHESIVE TABLET

Paris, April 9, 2009 – BioAlliance Pharma SA (Euronext Paris: BIO), the specialty pharmaceutical company focused on the treatment of opportunistic infections in cancer and AIDS, today announced that the FDA did not accept the NDA for Loramyc[®] (miconazole) mucoadhesive buccal tablet (MBT) to be filed based on the lack of a tablet imprint code. Loramyc[®] was approved in Europe in 2007 and is currently marketed in several EU territories including France, Germany, the UK, Sweden, Finland and Denmark. While the EU does not require a unique tablet identifier, the U.S. FDA does require a tablet imprint code for drug identification purposes. Prior to the initial filing, BioAlliance initiated the development of a debossed tablet to fulfill this requirement. BioAlliance will work closely with the FDA on the introduction of the debossed tablet and will soon after resubmit the Loramyc[®] application.

About BioAlliance Pharma

As a preferred partner for hospital-based specialists, BioAlliance Pharma is a specialty biopharmaceutical company which develops and markets innovative products, especially in the fields of opportunistic infections and chemotherapy complications. In areas where medical needs are insufficiently met, our targeted approaches help overcome drug resistance and improve patient health & quality of life. BioAlliance Pharma's ambition is to become a leading player in these fields by coupling innovation to patient needs. For more information, visit the BioAlliance Pharma web site at <http://www.bioalliancepharma.com>.

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This communication expressly or implicitly contains certain forward-looking statements concerning BioAlliance Pharma SA and its business. Such statements involve certain known and unknown risks, uncertainties and other factors, which could cause the actual results, financial condition, performance or achievements of BioAlliance Pharma SA to be materially different from any future results, performance or achievements expressed or implied by such forward-looking statements.

BioAlliance Pharma SA is providing this communication as of this date and does not undertake to update any forwardlooking statements contained herein as a result of new information, future events or otherwise. For a discussion of risks and uncertainties which could cause actual results, financial condition, performance or achievements of

BioAlliance Pharma SA to differ from those contained in the forward-looking statements, please refer to the Risk Factors ("Facteurs de Risque") section of the reference document approved by the AMF on April 11 2008 under the number R. 08-021, which is available on the AMF website (<http://www.amf-france.org>) or on BioAlliance Pharma S.A.'s website (<http://www.bioalliancepharma.com>).

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