



BioAlliance Pharma announces the grant of US patent covering Sitavig[®]

Paris, July 26, 2012 - BioAlliance Pharma SA (Euronext Paris - BIO), an innovative Company dedicated to the development of orphan oncology products and to supportive care products, announces the grant of a US patent protecting its product Sitavig[®]. This patent covers the Sitavig[®] tablet, its manufacturing process and its application for the treatment of recurrent labial herpes until 2027.

BioAlliance Pharma has conceived and developed Sitavig[®] (mucoadhesive tablet containing acyclovir) for the treatment of labial herpes in immunocompetent patients presenting more than 4 episodes a year. Sitavig[®] is an innovative mucoadhesive tablet to apply to the upper gum, allowing the delivery of very high concentrations of the active ingredient at the site of herpes infection, thus with a reinforced profile of efficacy.

"After the grant of a patent for Sitavig[®] in Europe and in major Asian territories, this US grant gives BioAlliance a worldwide protection for its product", declares Aude Michel, Vice President, Licensing and Legal Affairs, and European Patent Attorney of BioAlliance Pharma.

"We have obtained the receivability of Sitavig[®] US registration file last May from the FDA (Food and Drug Administration). This strong US patent protection allows its commercialization in the best conditions, reinforcing the value of this asset", added Judith Greciet, CEO of BioAlliance Pharma. "After a first license agreement with Teva for Israel and with a sales potential from \$120 to \$150 million, this product is becoming a suitable candidate to a license agreement, particularly in the United States".

About BioAlliance Pharma

Dedicated to cancer and supportive care treatment with a focus on resistance targeting and orphan products, BioAlliance conceives and develops innovative products, for specialty markets especially in the hospital setting and for orphan or rare diseases.

Created in 1997 and introduced to the Euronext Paris market in 2005, BioAlliance Pharma's ambition is to become a leading player in these fields by coupling innovation to patient needs. The company's teams have the key competencies required to identify, develop and register drugs in Europe and the USA.

BioAlliance Pharma has developed an advanced product portfolio:

Specialty products

Loramyc[®]/Oravig[®] (oropharyngeal candidiasis in immunocompromised patients): Registered in 28 countries (EU, US, Korea)

Sitavir[®]/Sitavig[®] (Acyclovir Lauriad[™]) (labialis herpes): Positive phase III final results; registration status

Fentanyl LauriadTM (chronic cancer pain): Positive preliminary Phase I results

Oncology Orphan products

Livatag[®] (Doxorubicin TransdrugTM) (primary liver cancer): Phase III on going

Validive[®] (Clonidine LauriadTM) (mucositis): Phase II on going

AMEP[®] (invasive melanoma): Phase I on going

For more information, visit the BioAlliance Pharma web site at www.bioalliancepharma.com

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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 2011 Reference Document filed with the AMF on April 24, 2012, which is available on the AMF website (<http://www.amf-france.org>) or on BioAlliance Pharma SA's website (<http://www.bioalliancepharma.com>).

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