UNITED STATES SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER PURSUANT TO RULE 13a-16 OR 15d-16 UNDER THE SECURITIES EXCHANGE ACT OF 1934

For the month of April 2016.

Commission File Number: 001-37384

GALAPAGOS NV

(Translation of registrant's name into English)

Generaal De Wittelaan L11 A3 2800 Mechelen, Belgium

(Address of principal executive office)
Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F. Form 20-F [x] Form 40-F []
Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):
Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):
On April 7, 2016 the Registrant issued a press release, a copy of which is attached hereto as Exhibit 99.1 and is incorporated herein by reference.
(c) Exhibit 99.1. Press release dated April 7, 2016
SIGNATURES
Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.
GALAPAGOS NV (Registrant)

/s/ XAVIER MAES

Xavier Maes

Company Secretary

Date: April 7, 2016

Galapagos and MorphoSys initiate Phase 1 study in joint antibody program MOR106

Mechelen, Belgium and Martinsried/Munich, Germany; 16.00 CET, 7 April 2016 - Galapagos NV (Euronext & NASDAQ: GLPG) and MorphoSys AG (FSE: MOR; Prime Standard Segment, TecDAX; OTC: MPSYY) announced today that dosing of MOR106, a first-in-class candidate human monoclonal antibody, has been initiated in healthy volunteers in a Phase 1 study. MOR106 was jointly discovered by Galapagos and MorphoSys under their collaboration, and has a novel mode of action with potential application in inflammation.

The primary objective of the Phase 1 study is to evaluate the safety and tolerability of single doses of MOR106. The study is randomized, double-blind, placebo-controlled and conducted in a single center in at least 56 healthy volunteers in Belgium, evaluating single ascending doses (SAD) administered as intravenous infusion. This Phase 1 study is characterized by its adaptive design, which enables the initiation of a subsequent multiple ascending dose (MAD) study in patients, depending on the outcome of the SAD study in healthy volunteers.

The study's secondary objective is to characterize the pharmacokinetic profile of MOR106 as well as monitor the occurrence of anti-drug antibodies as a measure of immunogenicity with MOR106. Topline results of the complete study, including the potential subsequent MAD part in patients, are expected in the second half of 2017.

"The alliance with MorphoSys has produced a first-in-class candidate human monoclonal antibody," said Piet Wigerinck, CSO of Galapagos. "MOR106 is the 10th Galapagos candidate with a new mode of action to be brought into the clinic."

"We are delighted to see the first antibody program from our alliance with Galapagos entering the clinical development stage. MOR106 is MorphoSys's fifth proprietary antibody program in clinical development and the first from our novel Ylanthia technology platform. We are excited about the growing value and maturity of our development pipeline. The start of this very innovative clinical development program also reflects the high value of our ongoing collaboration with Galapagos", commented Dr. Arndt Schottelius, Chief Development Officer of MorphoSys AG.

About MOR106 and the antibody collaboration

MOR106 is a human IgG1 monoclonal antibody for treatment of inflammatory diseases. MOR106 arises from the alliance initiated by Galapagos and MorphoSys in 2008, in which both companies contribute their core technologies and expertise. Galapagos provides the disease-related biology including cellular assays and targets discovered using its target discovery platform. MorphoSys contributes its Ylanthia antibody technology to generate fully human antibodies directed against the target and contributes full CMC development of this compound. Galapagos and MorphoSys will continue to equally share the research and development costs, as well as all future revenues.

About MorphoSys

MorphoSys developed HuCAL, the most successful antibody library technology in the pharmaceutical industry. By successfully applying this and other patented technologies, MorphoSys has become a leader in the field of therapeutic antibodies, one of the fastest-growing drug classes in human healthcare. Together with its pharmaceutical partners, MorphoSys has built a therapeutic pipeline of more than 100 human antibody drug candidates for the treatment of cancer, rheumatoid arthritis, and Alzheimer's disease, to name just a few. With its ongoing commitment to new antibody technology and drug development, MorphoSys is focused on making the healthcare products of tomorrow. MorphoSys is listed on the Frankfurt Stock Exchange under the symbol MOR. For regular updates about MorphoSys, visit http://www.morphosys.com.

HuCAL®, HuCAL GOLD®, HuCAL PLATINUM®, CysDisplay®, RapMAT®, arYla®, Ylanthia®, 100 billion high potentials®, Slonomics®, Lanthio Pharma® and LanthioPep® are registered trademarks of the MorphoSys Group.

About Galapagos

Galapagos(Euronext & NASDAQ: GLPG) is a clinical-stage biotechnology company specialized in the discovery and development of small molecule medicines with novel modes of action. Our maturing pipeline comprises Phase 2, Phase 1, pre-clinical, and discovery programs in cystic fibrosis, inflammation, fibrosis, osteoarthritis and other indications. We have discovered and developed filgotinib: in collaboration with Gilead we aim to bring this JAK1-selective inhibitor for inflammatory indications to patients all over the world. Galapagos is focused on the development and commercialization of novel medicines that will improve people's lives. The Galapagos group, including fee-for-service subsidiary Fidelta, has approximately 440 employees, operating from its Mechelen, Belgium headquarters and facilities in The Netherlands, France, and Croatia. More information at www.glpg.com.

Contacts

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Galapagos forward-looking statements

This release may contain forward-looking statements, including, among other things, statements regarding the mechanism of action and profile, and timing of clinical trials, of MOR106. Galapagos cautions the reader that forward-looking statements are not guarantees of future performance. Forward-looking statements involve known and unknown risks, uncertainties and other factors which might cause the actual results, financial condition and liquidity, performance or achievements of Galapagos, or industry results, to be materially different from any historic or future results, financial conditions and liquidity, performance or achievements expressed or implied by such forward-looking statements. In addition, even if Galapagos' results, performance, financial condition and liquidity, and the development of the industry in which it operates are consistent with such forward-looking statements, they may not be predictive of results or developments in future periods. Among the factors that may result in differences are that Galapagos' expectations regarding its MOR106 development program may be incorrect, the inherent uncertainties associated with competitive developments, clinical trial and product development activities and regulatory approval requirements (including that data from Galapagos' ongoing clinical research program may not support registration or further development of MOR106 due to safety, efficacy or other reasons), Galapagos' reliance on collaborations with third parties (including its collaboration partner for MOR106, MorphoSys), and estimating the commercial potential of MOR106. A further list and description of these risks, uncertainties and other risks can be found in Galapagos' securities and Exchange Commission (SEC) filings and reports, including in Galapagos' most recent annual report on form 20-F filed with the SEC and other filings and reports filed by Galapagos with the SEC. Given these uncertainties, the reader is advised not to place any undue reliance on such forward-looking statements. The