
**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**

April 2018

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 [] 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):

Note: Regulation S-T Rule 101(b)(1) only permits the submission in paper of a Form 6-K if submitted solely to provide an attached annual report to security holders.

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):

Note: Regulation S-T Rule 101(b)(7) only permits the submission in paper of a Form 6-K if submitted to furnish a report or other document that the registrant foreign private issuer must furnish and make public under the laws of the jurisdiction in which the registrant is incorporated, domiciled or legally organized (the registrant's "home country"), or under the rules of the home country exchange on which the registrant's securities are traded, as long as the report or other document is not a press release, is not required to be and has not been distributed to the registrant's security holders, and, if discussing a material event, has already been the subject of a Form 6-K submission or other Commission filing on EDGAR.

The information contained in this report on Form 6-K, including the Exhibit 99.1, except for the quote of Dr. Piet Wigerinck contained in Exhibit 99.1, is hereby incorporated by reference into the Company's Registration Statements on Forms F-3 (File No. 333-211765) and S-8 (File Nos. 333-204567, 333-208697, 333-211834, 333-215783, and 333-218160).

On April 24, 2018, 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 24, 2018

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)

Date: April 24, 2018

/s/ Xavier Maes

Xavier Maes

Company Secretary

Galapagos reports initiation of FALCON clinical trial in cystic fibrosis

- **Our first triple combination investigational therapy for CF patients**
 - combination of '2451+'2222+'2737
 - multiple centers in Europe
 - topline results expected Q3 2018

Mechelen, Belgium; 24 April 2018; 08.45 AM CET - Galapagos NV (Euronext & NASDAQ: GLPG) announces initiation of its first clinical trial with an investigational triple combination therapy in cystic fibrosis patients.

The aim of the FALCON trial is to evaluate the efficacy, safety, tolerability, and pharmacokinetics of a novel triple combination in up to 24 CF patients. The open label trial is being conducted in multiple centers, initially in the United Kingdom with expansion expected to other European countries.

The FALCON study will comprise two parts. Part one will entail treatment of 8 patients for two weeks with a fixed dose dual combination of potentiator GLPG2451 and C1 corrector GLPG2222 in homozygous F508del patients. This will be followed by two weeks' treatment with '2451, '2222, and C2 corrector GLPG2737. Part two will entail treatment for two weeks with a higher dose dual combination of '2451 and '2222 in separate 8 patient cohorts of homozygous F508del patients and heterozygous F508del patients with a minimal function mutation on the other allele. This will be followed by two weeks' treatment with '2451, '2222, and '2737. Efficacy will be measured by changes in sweat chloride and percent predicted forced expiratory volume during the first second (ppFEV1%). Topline results from treatment in part one of FALCON are expected to be disclosed Q3 2018.

"Today marks a key milestone in our CF program. Since 2005, it has been our ambition to develop a disease-modifying therapy for CF. Now we are evaluating a first triple combination in CF patients, with the next triple combination coming up rapidly behind it," said Dr. Piet Wigerinck, CSO of Galapagos. "These steps bring us closer to our goal of offering patients, doctors, and payers more choice in their CF therapies."

GLPG2451, GLPG2222, GLPG2737 (and combinations thereof) are investigational therapies; their safety and efficacy have not been established.

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. Galapagos' pipeline comprises Phase 3 through to discovery programs in cystic fibrosis, inflammation, fibrosis, osteoarthritis and other indications. Our target discovery platform has delivered three novel mechanisms showing promising patient results in, respectively, inflammatory diseases, idiopathic pulmonary fibrosis and atopic dermatitis. 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 600 employees, operating from its Mechelen, Belgium headquarters and facilities in the Netherlands, France, Switzerland, the US and Croatia. More information at www.glp.com.

Contacts

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Forward-looking statements

This release may contain forward-looking statements, including statements regarding Galapagos' strategic ambitions, the composition of a potential triple combination therapy for CF; the potential activity of GLPG2451, GLPG2222, GLPG2737 and/or of combinations thereof in CF; the anticipated timing of clinical trials with, and plans related to, the CF portfolio of product candidates; the timing, progression and/or results of such trials and plans. 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 the inherent uncertainties associated with competitive developments, clinical trial and product development activities and regulatory approval requirements (including that data from the ongoing and planned clinical research programs in CF may not support registration or further development of a potential triple combination or any of Galapagos' potentiators or correctors (alone or in combination) due to safety, efficacy or other reasons), Galapagos' reliance on collaborations with third parties (including its collaboration partner for CF, AbbVie), and estimating the commercial potential of Galapagos' CF portfolio. 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 subsequent 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. These forward-looking statements speak only as of the date of publication of this document. Galapagos expressly disclaims any obligation to update any such forward-looking statements in this document to reflect any change in its expectations with regard thereto or any change in events, conditions or circumstances on which any such statement is based or that may affect the likelihood that actual results will differ from those set forth in the forward-looking statements, unless specifically required by law or regulation.