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 November 2020

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):

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 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. 333-230639) and S-8 (File Nos. 333-204567, 333-208697, 333-211834, 333-215783, 333-218160, 333-225263, 333-231765 and 333-249416).

On November 9, 2020, 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 November 9, 2020

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: November 9, 2020

/s/ Xavier Maes
Xavier Maes
Company Secretary

First patient dosed with GLPG3667 in psoriasis patient Phase 1b trial

Mechelen, Belgium; 9 November 2020, 22.01 CET – Galapagos NV (Euronext & NASDAQ: GLPG) announces dosing of the first psoriasis patient in the Phase 1b trial with GLPG3667.

GLPG3667 is a proprietary compound with an undisclosed mechanism of action in development for inflammatory and autoimmune indications. Based on an initial clinical study in healthy volunteers, GLPG3667 has met the criteria to be further studied in patients.

This Phase 1b trial is a randomized, double-blind, placebo-controlled, multicenter study to evaluate the safety, tolerability, efficacy, pharmacokinetics, and pharmacodynamics of GLPG3667. A daily oral administration of GLPG3667 at two different dose levels or a placebo will be investigated for 4 weeks in 30 patients with moderate to severe plaque psoriasis. The primary endpoint will be the change from baseline in Psoriasis Area Severity Index (PASI) score at 4 weeks. The first patient was dosed today. Recruitment will be based in Europe.

Pending successful completion of the Phase 1b study in psoriasis, we anticipate evaluating GLPG3667 in dose range finding Phase 2 studies in psoriatic arthritis and ulcerative colitis in the second half of 2021.

“We continue to press forward with several mechanisms of action in inflammation as part of our strategy to bring improved therapies to patients. We look forward to applying our extensive clinical experience in inflammation to this additional approach with GLPG3667. The study in psoriasis and the biomarker information collected should pave the way for trials in psoriatic arthritis and other indications,” said Dr. Piet Wigerinck, Chief Scientific Officer of Galapagos.

GLPG3667 is an investigational drug and not approved by any regulatory authority. Its efficacy and safety have not been established.

About our early inflammation pipeline
Galapagos continues to innovate in the inflammation space, with multiple, distinct mechanism of action approaches to bringing new therapy options to patients. GLPG3667 is an oral small molecule with undisclosed mechanism of action currently in Phase 1 trials.

For more information about our early clinical programs: www.glpg.com/other-programs
For information about the studies with GLPG3667 in psoriasis (NCT04594928): www.clinicaltrials.gov

About Galapagos
Galapagos NV discovers and develops small molecule medicines with novel modes of action, several of which show promising patient results and are currently in late-stage development in multiple diseases. Our pipeline comprises discovery through Phase 3 programs in inflammation, fibrosis and other indications. Our ambition is to become a leading global biopharmaceutical company focused on the discovery, development and commercialization of innovative medicines. More information at www.glpg.com.

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Forward-looking statements

This press release includes forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, that are subject to risks, uncertainties and other factors that could cause actual results to differ materially from those referred to in the forward-looking statements and, therefore, the reader should not place undue reliance on them. These risks, uncertainties and other factors include, without limitation, the risk that ongoing and future clinical studies with

GLPG3667 may not be completed in the currently envisaged timelines or at all, 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 may not support registration or further development of GLPG3667 due to safety, efficacy or other reasons), that Galapagos' may not be able to fulfil its strategic ambition to bring therapies with different mechanism of action to patients with inflammatory diseases, Galapagos' reliance on collaborations with third parties and that Galapagos' estimations regarding its GLPG3667 development program and regarding the commercial potential of GLPG3667, may be incorrect, as well as those risks and uncertainties identified in our Annual Report on Form 20-F for the year ended 31 December 2019 and our subsequent filings with the SEC. All statements other than statements of historical fact are statements that could be deemed forward-looking statements. The forward-looking statements contained herein are based on management's current expectations and beliefs and speak only as of the date hereof, and Galapagos makes no commitment to update or publicly release any revisions to forward-looking statements in order to reflect new information or subsequent events, circumstances or changes in expectations.