
**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 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 and the quote of Mr. Pawel Przewiezlikowski 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 and 333-231765).

On April 16, 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 April 16, 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: April 16, 2020

/s/ Xavier Maes

Xavier Maes
Company Secretary

Galapagos and Ryvu announce research collaboration

Mechelen, Belgium and Krakow, Poland; 16 April 2020, 07.30 CET – Galapagos NV (Euronext & NASDAQ: GLPG) and Ryvu Therapeutics S.A. (WSE: RVU) today announced a collaboration focused on the discovery and development of novel small molecule drugs in inflammation.

Ryvu specializes in the discovery and development of first-in-class small molecules and drug candidates in diseases with high unmet medical needs. The collaboration announced today is based on a novel drug target identified by Ryvu, which will contribute its technology platform and related intellectual property (IP). Ryvu and Galapagos will both provide resources to support the collaboration and make use of their expertise in high-throughput screening, biology, medicinal chemistry, and toxicology.

This is a joint research collaboration in which Ryvu is responsible for early drug discovery. Under the terms of the agreement, Galapagos will have an exclusive option to license IP developed by Ryvu and to continue to develop this during the collaboration. Pending achievement of pre-agreed criteria and utilizing its option, Galapagos will be responsible for all further development of the program.

In exchange for global development and commercialization rights, Ryvu will receive an upfront payment and will be eligible for further option, milestone, and royalty payments.

“We believe that the collaboration with Ryvu is an excellent fit, as both companies are driven by the search for novel drugs to address unmet medical needs,” says Dr. Piet Wigerinck, Chief Scientific Officer at Galapagos. “We look forward to collaborating with the Ryvu team to push this program forward.”

Pawel Przewiezlikowski, Ryvu Chief Executive Officer, added: “We are thrilled to start working with Galapagos, a real role model for the European biotechnology sector. Throughout joint discussions on the collaboration, our teams have developed a strong rapport and built the foundations for a very promising start for the new project on an exciting novel target.”

About Galapagos

Galapagos (Euronext & NASDAQ: GLPG) discovers and develops small molecule medicines with novel modes of action, three 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, osteoarthritis 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.glp.com.

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Forward-looking statements

This release may contain forward-looking statements, including, among other things, statements regarding the success of the collaboration with Ryvu, the identification and validation of a target by Ryvu, the potential future exercise of any option granted to Galapagos, the mechanism of action and profile of, and timing and results of clinical trials with, and potential commercialization of, any compounds coming out of any in-licensed program, as well as statements regarding potential future option exercise, milestone and royalty payments. 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 the further development of any potential future in-licensed program, including its potential to address a large unmet need in inflammation, may be incorrect, the inherent uncertainties associated with competitive developments, clinical trial and product development activities and regulatory approval requirements (including that data from research and development programs may not support further development of the compound(s) due to safety, efficacy or other reasons), Galapagos' reliance on collaborations with third parties and estimating the commercial potential of Galapagos' product candidates. 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. 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.

About Ryvu Therapeutics

Ryvu Therapeutics is a clinical stage biopharmaceutical company developing novel small molecule therapies that address emerging targets in oncology. Pipeline candidates make use of diverse therapeutic mechanisms driven by emerging knowledge of cancer biology, including small molecules directed at kinase, synthetic lethality, immuno-oncology and cancer metabolism targets. SEL120 is a selective CDK8 kinase inhibitor with potential for the treatment of hematological malignancies and solid tumors currently in clinical development for the treatment of acute myeloid leukemia and myelodysplastic syndrome. SEL24/MEN1703 is a dual PIM/FLT3 kinase inhibitor licensed to the Menarini Group in clinical development for the treatment of acute myeloid leukemia. Ryvu is listed on the Warsaw Stock Exchange in Poland (WSE:RVU). For more information, please see www.ryvu.com.

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