UNITED STATES SECURITIES AND EXCHANGE COMMISSION

| Washington, D.C. 20549 |
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| 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 May 2023 |
| Commission File Number: 001-41421 |
| Alvotech (Translation of registrant's name into English) |
| (Hansiation of registrant's name into Engusit) |
| 9, Rue de Bitbourg, L-1273 Luxembourg, Grand Duchy of Luxembourg (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 |

INFORMATION CONTAINED IN THIS REPORT ON FORM 6-K

Incorporation by reference

This Report on Form 6-K (this "Report") of Alvotech (the "Company"), including Exhibit 99.1 attached hereto, is being furnished and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934 (the "Exchange Act") or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933 or the Exchange Act.

Press Release

On May 4, 2023, Alvotech announced the initiation of a confirmatory patient study for AVT05, a proposed biosimilar for Simponi[®] and Simponi Aria[®] (golimumab). A copy of the announcement is furnished as Exhibit 99.1 to this Report on Form 6-K.

EXHIBIT INDEX

Exhibit No.

No. Description

99.1 <u>Press Release dated May 4, 2023.</u>

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.

Date: May 4, 2023

ALVOTECH

By: /s/ Tanya Zharov

Name: Tanya Zharov Title: General Counsel



Alvotech Initiates Confirmatory Patient Study for AVT05, a Proposed Biosimilar for Simponi® and Simponi Aria®

• The clinical study will compare AVT05 and Simponi® in patients with moderate to severe rheumatoid arthritis

REYKJAVIK, ICELAND (May 4, 2023) – Alvotech (NASDAQ: ALVO), a global biotech company specializing in the development and manufacture of biosimilar medicines for patients worldwide, announced today the initiation of Alvotech's confirmatory patient study for AVT05, a biosimilar candidate to Simponi[®] and Simponi Aria[®] (golimumab). The objective of the clinical study is to compare the efficacy, safety, and immunogenicity of AVT05 and Simponi[®] in adult patients with moderate to severe rheumatoid arthritis.

"It is a pleasure to be able to mark another clinical milestone in our biosimilars development pipeline," said Joseph McClellan, Chief Scientific Officer. "Alvotech's platform approach allows us to diversify and develop multiple product candidates in parallel. This is part of our strong commitment to improve patient lives by expanding access to affordable biologics globally."

The AVT05 confirmatory patient study (https://clinicaltrials.gov/ct2/show/NCT05842213) is a randomized, double-blind, 2-arm multicenter study comparing the efficacy, safety, and immunogenicity of subcutaneous AVT05 and EU-approved Simponi in patients with moderate to severe rheumatoid arthritis. The primary outcome measure will be to demonstrate comparative efficacy of AVT05 with Simponi at week 16 based on change from baseline on a standardized Disease Activity Standard score (DAS28-CRP) which assesses the severity of rheumatoid arthritis using clinical and laboratory data. In January 2023 Alvotech announced the initiation of a pharmacokinetic study for AVT05.

In the twelve months up to April 2023, combined net revenues worldwide from sales of Simponi and Simponi Aria were over \$2.1 billion, based on reported sales by the manufacturer of the reference product [1].

[1] Source: Johnson & Johnson quarterly reports.

About AVT05

AVT05 is a biosimilar candidate for Simponi and Simponi Aria (golimumab). Golimumab is a monoclonal antibody that inhibits tumor necrosis factor alpha (TNF alpha), a cytokine protein in the body. Elevated TNF alpha levels have been implicated in several chronic inflammatory diseases such as rheumatoid arthritis, psoriatic arthritis, and ankylosing spondylitis. AVT05 is an investigational product and has not received regulatory approval in any country. Biosimilarity has not been established by regulatory authorities and is not claimed.

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About Alvotech

Alvotech is a biotech company, founded by Robert Wessman, focused solely on the development and manufacture of biosimilar medicines for patients worldwide. Alvotech seeks to be a global leader in the biosimilar space by delivering high quality, cost-effective products, and services, enabled by a fully integrated approach and broad in-house capabilities. Alvotech's current pipeline contains eight biosimilar candidates aimed at treating autoimmune disorders, eye disorders, osteoporosis, respiratory disease, and cancer. Alvotech has formed a network of strategic commercial partnerships to provide global reach and leverage local expertise in markets that include the United States, Europe, Japan, China, and other Asian countries and large parts of South America, Africa and the Middle East. Alvotech's commercial partners include Teva Pharmaceuticals, a US affiliate of Teva Pharmaceutical Industries Ltd. (US), STADA Arzneimittel AG (EU and select other territories), Fuji Pharma Co., Ltd (Japan), Advanz Pharma (EEA, UK, Switzerland, Canada, Australia and New Zealand), Cipla/Cipla Gulf/Cipla Med Pro (Australia, New Zealand, South Africa), JAMP Pharma Corporation (Canada), Yangtze River Pharmaceutical (Group) Co., Ltd. (China), DKSH (Taiwan, Hong Kong, Cambodia, Malaysia, Singapore, Indonesia, India, Bangladesh and Pakistan), YAS Holding LLC (Middle East and North Africa), Abdi Ibrahim (Turkey), Kamada Ltd. (Israel), Mega Labs, Stein, Libbs, Tuteur and Saval (Latin America) and Lotus Pharmaceuticals Co., Ltd. (Thailand, Vietnam, Philippines, and South Korea). Each commercial partnership covers a unique set of product(s) and territories. Except as specifically set forth therein, Alvotech disclaims responsibility for the content of periodic filings, disclosures and other reports made available by its partners. For more information, please visit www.alvotech.com. None of the information on the Alvotech website shall be deemed part of this press release.

Forward-Looking Statements

Certain statements in this communication may be considered "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, as amended. Forward-looking statements generally relate to future events or the future financial operating performance of Alvotech and may include, for example, Alvotech's expectations regarding competitive advantages, business prospects and opportunities including pipeline product development, future plans and intentions, results, level of activities, performance, goals or achievements or other future events, regulatory review and interactions, the satisfactory responses to the FDA's inspection findings and resolution of other deficiencies conveyed following the inspection of Alvotech's manufacturing site, the potential approval and commercial launch of its product candidates, the timing of regulatory approval and market launches, and the estimated size of the total addressable market of Alvotech's pipeline products. In some cases, you can identify forward-looking statements by terminology such as "may", "should", "expect", "intend", "will", "estimate", "anticipate", "believe", "predict", "potential", "aim" or "continue", or the negatives of these terms or variations of them or similar terminology. Such forward-looking statements are subject to risks, uncertainties, and other factors which could cause actual results to differ materially from those expressed or implied by such forward-looking statements. These forward-looking statements are based upon estimates and assumptions that, while considered reasonable by Alvotech and its management, are inherently uncertain and are inherently subject to risks, variability, and contingencies, many of which are beyond Alvotech's control. Factors that may cause actual results to differ materially from current expectations include, but are not limited to: (1) the outcome of any

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legal proceedings that may be instituted against Alvotech or others following the business combination between Alvotech Holdings S.A., Oaktree Acquisition Corp. II and Alvotech: (2) the ability to maintain stock exchange listing standards: (3) changes in applicable laws or regulations: (4) the possibility that Alvotech may be adversely affected by other economic, business, and/or competitive factors; (5) Alvotech's estimates of expenses and profitability; (6) Alvotech's ability to develop, manufacture and commercialize the products and product candidates in its pipeline; (7) actions of regulatory authorities, which may affect the initiation, timing and progress of clinical studies or future regulatory approvals or marketing authorizations; (8) the ability of Alvotech or its partners to respond to inspection findings and resolve deficiencies to the satisfaction of the regulators; (9) the ability of Alvotech or its partners to enroll and retain patients in clinical studies; (10) the ability of Alvotech or its partners to gain approval from regulators for planned clinical studies, study plans or sites; (11) the ability of Alvotech's partners to conduct, supervise and monitor existing and potential future clinical studies, which may impact development timelines and plans; (12) Alvotech's ability to obtain and maintain regulatory approval or authorizations of its products, including the timing or likelihood of expansion into additional markets or geographies; (13) the success of Alvotech's current and future collaborations, joint ventures, partnerships or licensing arrangements; (14) Alvotech's ability, and that of its commercial partners, to execute their commercialization strategy for approved products; (15) Alvotech's ability to manufacture sufficient commercial supply of its approved products; (16) the outcome of ongoing and future litigation regarding Alvotech's products and product candidates; (17) the potential impact of the ongoing COVID-19 pandemic on the FDA's review timelines, including its ability to complete timely inspection of manufacturing sites; (18) the impact of worsening macroeconomic conditions, including rising inflation and interest rates and general market conditions, war in Ukraine and global geopolitical tension, and the ongoing and evolving COVID-19 pandemic on the Company's business, financial position, strategy and anticipated milestones; and (19) other risks and uncertainties set forth in the sections entitled "Risk Factors" and "Cautionary Note Regarding Forward-Looking Statements" in documents that Alvotech may from time to time file or furnish with the SEC. There may be additional risks that Alvotech does not presently know or that Alvotech currently believes are immaterial that could also cause actual results to differ from those contained in the forward-looking statements. Nothing in this communication should be regarded as a representation by any person that the forward-looking statements set forth herein will be achieved or that any of the contemplated results of such forward-looking statements will be achieved. You should not place undue reliance on forwardlooking statements, which speak only as of the date they are made. Alvotech does not undertake any duty to update these forward-looking statements or to inform the recipient of any matters of which any of them becomes aware of which may affect any matter referred to in this communication. Alvotech disclaims any and all liability for any loss or damage (whether foreseeable or not) suffered or incurred by any person or entity as a result of anything contained or omitted from this communication and such liability is expressly disclaimed. The recipient agrees that it shall not seek to sue or otherwise hold Alvotech or any of its directors, officers, employees, affiliates, agents, advisors, or representatives liable in any respect for the provision of this communication, the information contained in this communication, or the omission of any information from this communication.

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