
**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 December 2025

Commission File Number: **001-41421**

Alvotech

(Translation of registrant's name into English)

**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 [X] Form 40-F []

Incorporation by Reference

This Report on Form 6-K (this “Report”) of Alvotech (the “Company”) excluding Exhibit 99.1 attached hereto, shall be deemed to be incorporated by reference into the Company’s registration statements on Forms F-3 (File Nos. 333-266136, 333-273262, 333-275111 and 333-281684), the Company’s registration statement on Form F-3ASR (File No. 333-289006), and the Company’s registration statement on Form S-8 (File No. 333-266881) and to be a part thereof from the date on which this Report is filed, to the extent not superseded by documents or reports subsequently filed or furnished. Exhibit 99.1 to this Report 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 Releases

On December 22, 2025, Alvotech issued a Press Release announcing that launches have commenced in Europe for Gobivaz® (golimumab), Alvotech’s biosimilar to Simponi® (golimumab), also known as AVT05 (prefilled syringe and autoinjector). A copy of the Press Release is furnished herewith as exhibit 99.1.

EXHIBIT INDEX

<u>Exhibit Number</u>	<u>Description</u>
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99.1	Press Release dated December 22, 2025
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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.

Alvotech
(Registrant)

Date: December 22, 2025

/s/ Tanya Zharov
Tanya Zharov
General Counsel

Alvotech announces European launch of first-in-market biosimilar to Simponi® (golimumab) globally by partner Advanz Pharma

- *Introduction of AVT05 supported by NHS England tender award in the United Kingdom*

REYKJAVIK, Iceland, Dec. 22, 2025 (GLOBE NEWSWIRE) -- Alvotech (NASDAQ: ALVO), a global biotechnology company specializing in the development and manufacture of biosimilar medicines for patients worldwide, today announced that launches are underway in Europe for Gobivaz® (golimumab), Alvotech's biosimilar to Simponi® (golimumab), also known as AVT05 (prefilled syringe and autoinjector). Gobivaz is the first marketed biosimilar to Simponi worldwide. It is being commercialized exclusively in Europe by Alvotech's partner, Advanz Pharma Holdco Limited ("Advanz Pharma").

In the United Kingdom, the introduction of Gobivaz is supported by a National Health Service (NHS) England tender award. This procurement decision provides a defined framework for adoption within NHS services and reflects the system's focus on improving access to high quality biologic treatment options for immune mediated inflammatory diseases.

Availability across EEA markets will progress in line with national pricing and reimbursement processes.

"We welcome the introduction of Gobivaz into clinical practice across Europe following its approval as the first biosimilar to Simponi worldwide," said Robert Wessman, Chairman and Chief Executive Officer of Alvotech. "Our biosimilar represents an important treatment option for patients, and the NHS England tender supports our aim of expanding the availability of high-quality biologic medicines."

Approval of Gobivaz was supported by comprehensive analytical, pre-clinical and clinical studies that demonstrated biosimilarity to the reference product. Gobivaz is available in the same presentations and dosage forms as Simponi and is manufactured at Alvotech's facility in Iceland.

For further information, please contact:

Media

Benedikt Stefansson, VP Investor Relations and Global Communications
Sarah Macleod, Head of Global Communications
alvotech.media@alvotech.com

Investor Relations

Balaji Prasad, Chief Strategy Officer
Patrik Ling, VP Investor Relations Scandinavia (SE)
Benedikt Stefansson, VP Investor Relations and Global Communications (IS)
alvotech.ir@alvotech.com

About Alvotech

Alvotech is a biotechnology 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. Five biosimilars are already approved and marketed in multiple global markets, including biosimilars to Humira® (adalimumab), Stelara® (ustekinumab), Simponi® (golimumab), Eylea® (aflibercept) and Prolia®/Xgeva® (denosumab). The current development pipeline includes nine disclosed 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. For more information, please visit <https://www.alvotech.com>. None of the information on the Alvotech website shall be deemed part of this press release.

For more information, please visit our investor portal, and our website or follow us on social media on LinkedIn, Facebook, Instagram and YouTube.

Alvotech 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 include, for example, Alvotech's expectations regarding competitive advantages, business prospects and opportunities including pipeline product development, future plans and intentions, regulatory submissions, review and interactions, the potential approval and commercial launch of its product candidates, the timing of regulatory approval, market launches and financial projections. 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 factors 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. These forward-looking statements are provided for illustrative purposes only and are not intended to serve as, and must not be relied on by an investor as, a guarantee, assurance, prediction or definitive statement of a fact or probability. 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.