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 July 2026

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Form S-8 (File No. 333-266881) and to be a part thereof from the date on which this Report is filed, to the extent not superseded by documents or reports subsequently filed or furnished.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 Release

On July 29, 2026, Alvotech issued a Press Release announcing that the US Food and Drug Administration has closed its inspection of the company's manufacturing site, confirming the inspection classification of the site is Voluntary Action Indicated (VAI). A copy of the Press Release is furnished herewith as exhibit 99.1.

EXHIBIT INDEX

Exhibit Number	Description
99.1	Press Release dated July 29, 2026

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: July 29, 2026

/s/ Tanya Zharov
Tanya Zharov
General Counsel

EXHIBIT 99.1

Alvotech announces successful closure of FDA inspection of manufacturing facility

REYKJAVIK, Iceland, July 29, 2026 (GLOBE NEWSWIRE) -- Alvotech (NASDAQ: ALVO; ALVO-SDB) (the "Company"), a global biotechnology company specializing in the development and manufacture of biosimilar medicines for patients worldwide, today announced that the U.S. Food and Drug Administration (FDA) has closed its inspection of the company's manufacturing facility in Reykjavik, Iceland, conducted in May 2026. The FDA has confirmed the inspection classification of the manufacturing site is Voluntary Action Indicated (VAI).

"This is an important milestone for Alvotech and reflects the significant work our teams have undertaken to further strengthen our quality systems and manufacturing operations," said Lisa Graver, CEO of Alvotech. "With the conclusion of the FDA's recent inspection, we are confident that the actions we have taken have effectively addressed the observations identified. We continue to work with the FDA to advance our recent Biologics Licence Applications towards approval."

Alvotech continues to advance its pipeline of biosimilar candidates. The company recently announced the resubmission of Biologics License Applications (BLAs) for AVT05, Alvotech's biosimilar to Simponi® (golimumab), and AVT06, Alvotech's biosimilar to Eylea® (aflibercept), and the acceptance by the FDA of the submission of a BLA for AVT16, Alvotech's proposed biosimilar to Entyvio® (vedolizumab) to the FDA in June 2026.

Alvotech continues to work closely with health authorities worldwide to bring high-quality, affordable biologic medicines to patients.

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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 13 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.

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

None of the information on the Alvotech website shall be deemed part of this press release.

Use of trademarks

Simponi® and Simponi Aria® are trademarks of Johnson & Johnson. Eylea® is a trademark of Regeneron Pharmaceuticals Inc. Entyvio® is a registered trademark of Millennium Pharmaceuticals, Inc.