UNITED STATES INTERNATIONAL TRADE COMMISSION
Washington, D.C.

In the Matter of **CERTAIN ANTIBODY DRUG CONJUGATES AND COMPONENTS THEREOF AND PRODUCTS CONTAINING THE SAME**	**Investigation No. 337-TA-1466**

NOTICE OF A COMMISSION DETERMINATION NOT TO REVIEW AN INITIAL DETERMINATION SETTING THE TARGET DATE

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission ("Commission") has determined not to review an initial determination ("ID") (Order No. 5) of the presiding administrative law judge ("ALJ"), setting a 17-month target date. Accordingly, the target date for this investigation is May 21, 2027, and the final ID on violation shall issue by January 21, 2027.

FOR FURTHER INFORMATION CONTACT: Ronald A. Traud, Esq., Office of the General Counsel, U.S. International Trade Commission, 500 E Street S.W., Washington, D.C. 20436, telephone (202) 205-3427. Copies of non-confidential documents filed in connection with this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov*. For help accessing EDIS, please email EDIS3Help@usitc.gov. General information concerning the Commission may also be obtained by accessing its Internet server at *https://www.usitc.gov*. Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205-1810.

SUPPLEMENTARY INFORMATION: The Commission instituted this investigation on December 22, 2025, based on a complaint filed by AbbVie Inc. of North Chicago, Illinois; ImmunoGen, Inc. of Waltham, Massachusetts; and ImmunoGen Switzerland GmbH. 90 FR 59,867 (Dec. 22, 2025). The complaint, as supplemented, alleges violations of section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, in the importation into the United States of certain antibody drug conjugates and components thereof and products containing the same by reason of misappropriation of trade secrets, the threat or effect of which is to destroy or substantially injure an industry in the United States or to prevent the establishment of an industry in the United States. *Id*. The Commission's notice of investigation named as respondents ProfoundBio US Co. of Seattle, Washington; ProfoundBio (Suzhou) Co., Ltd. of Suzhou, China; Genmab A/S of Valby, Denmark; Genmab B.V. of Utrecht, Netherlands; and Genmab US, Inc.

of Plainsboro, New Jersey. *Id*. The Office of Unfair Import Investigations is participating in the investigation. *Id*.

On January 13, 2026, the ALJ issued Order No. 5 (the subject ID), setting a 17-month target date. According to the ID, the target date for this investigation is May 21, 2027, and the final ID on violation shall issue by January 21, 2027. No petitions for review of the ID were filed.

The Commission has determined not to review the subject ID. Accordingly, the target date is May 21, 2027, and the final ID on violation shall issue by January 21, 2027.

The Commission vote for this determination took place on February 9, 2026.

The authority for the Commission's determination is contained in section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in Part 210 of the Commission's Rules of Practice and Procedure (19 CFR Part 210).

By order of the Commission.

Lisa R. Barton
Secretary to the Commission

Issued: February 10, 2026