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2026/0075 (NLE)**

POLCOM 131

PROPOSAL

From:	Secretary-General of the European Commission, signed by Ms Martine DEPREZ, Director
date of receipt:	10 April 2026
To:	Ms Thérèse BLANCHET, Secretary-General of the Council of the European Union

No. Cion doc.:	COM(2026) 130 annex
Subject:	ANNEX to the Proposal for a Council Decision on the position to be taken on behalf of the European Union within the CETA Joint Committee established under the Comprehensive Economic and Trade Agreement (CETA) between Canada, of the one part, and the European Union and its Member States, of the other part, as regards the adoption of a decision concerning the inclusion of active pharmaceutical ingredients as the medicinal products or drugs listed in paragraph 2 of the Annex 1 to the Protocol on the mutual recognition of the compliance and enforcement programme regarding good manufacturing practices for pharmaceutical products

Delegations will find attached document COM(2026) 130 annex.

Encl.: COM(2026) 130 annex



Brussels, 10.4.2026
COM(2026) 130 final

ANNEX

ANNEX

to the

Proposal for a Council Decision

on the position to be taken on behalf of the European Union within the CETA Joint Committee established under the Comprehensive Economic and Trade Agreement (CETA) between Canada, of the one part, and the European Union and its Member States, of the other part, as regards the adoption of a decision concerning the inclusion of active pharmaceutical ingredients as the medicinal products or drugs listed in paragraph 2 of the Annex 1 to the Protocol on the mutual recognition of the compliance and enforcement programme regarding good manufacturing practices for pharmaceutical products

ANNEX

Draft

DECISION No.../2026 OF THE CETA JOINT COMMITTEE of [date]

as regards the inclusion of active pharmaceutical ingredients as the medicinal products or drugs listed in paragraph 2 of the Annex 1 to the Protocol on the mutual recognition of the compliance and enforcement programme regarding good manufacturing practices for pharmaceutical products

THE CETA JOINT COMMITTEE,

Having regard to Article 26.1 of the *Comprehensive Economic and Trade Agreement (CETA) between Canada, on the one part, and the European Union and its Member States, on the other part*, done at Brussels on 30 October 2016 (the 'Agreement'),

Whereas:

- (1) Under the provisions of Article 30.7.3(a), the Agreement has been provisionally applied since September 21, 2017.
- (2) Article 26.1.5(c) of the Agreement provides that the CETA Joint Committee may consider or agree on amendments as provided in the Agreement.
- (3) Article 30.2.2 of the Agreement provides that the CETA Joint Committee may decide to amend the protocols and annexes of the Agreement.
- (4) Article 15 paragraphs 5 and 6 of the *Protocol on the mutual recognition of the compliance and enforcement programme regarding good manufacturing practices for pharmaceutical products* (the 'Protocol') provides that the Joint Sectoral Group is to review the operational scope of medicinal products or drugs under paragraph 2 of Annex 1, with a view to including those medicinal products or drugs listed in paragraph 1 of Annex 1.
- (5) The European Commission and Health Canada have conducted evaluations of the applicable Canadian and EU Good Manufacturing Practices programs for active pharmaceutical ingredients and have concluded that their respective regulatory and enforcement frameworks in this area are equivalent.
- (6) The Protocol, on the recommendation of the Joint Sectoral Group to the CETA Joint Committee, should therefore be amended by including active pharmaceutical ingredients as medicinal products or drugs listed in paragraph 2 of Annex 1 to the Protocol.
- (7) The entry into force of this Decision does not require any further procedure under the European Union legal order.

HAS ADOPTED THIS DECISION:

Article 1

Operational scope of medicinal products or drugs

Paragraph 2 of Annex 1 to the Protocol on the mutual recognition of the compliance and enforcement programme regarding good manufacturing practices for pharmaceutical products shall be amended as follows:

- (a) at the end of the entry for point (f), the word “and” shall be removed;
- (b) at the end of the entry for point (g), the full stop shall be replaced by “; and”; and,
- (c) after the entry for point (g), the following entry shall be added: “(h) active pharmaceutical ingredients.”.

Article 2

Entry into force

This Decision shall enter into force on the first day of the second month following the date Canada has provided to the European Union a written notification certifying that it has completed its internal requirements and procedures necessary for entry into force.

Article 3

Legal status of the amendment

Upon entry into force of this Decision, the amendment set out in Article 1 will become part of the Protocol. The Parties recognize that since the Agreement, including the Protocol, is provisionally applied, the amendment will also be provisionally applied until the Agreement enters into force.

Article 4

Authentic texts

This Decision is drawn up in duplicate in the Bulgarian, Croatian, Czech, Danish, Dutch, English, Estonian, Finnish, French, German, Greek, Hungarian, Italian, Latvian, Lithuanian, Maltese, Polish, Portuguese, Romanian, Slovak, Slovenian, Spanish and Swedish languages, each version being equally authentic.

*For the Joint Committee
The Co-chairs*