NOTIFICATION

Addendum

The following communication, dated 4 October 2024, is being circulated at the request of the delegation of Ukraine.

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**Title:** Draft Resolution of the Cabinet of Ministers of Ukraine "Some Іssues of Safety and Verification of Medicinal Products"

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| **Reason for Addendum:** |
| [ ] | Comment period changed - date:  |
| [X] | Notified measure adopted - date: 26 September 2024 |
| [X] | Notified measure published - date: 3 October 2024 |
| [X] | Notified measure enters into force - date: 3 October 2024; |
| [X] | Text of final measure available from[[1]](#footnote-1): <https://zakon.rada.gov.ua/laws/show/1121-2024-%D0%BF#Text><https://members.wto.org/crnattachments/2024/TBT/UKR/final_measure/24_06523_00_x.pdf> |
| [ ] | Notified measure withdrawn or revoked - date: Relevant symbol if measure re-notified:  |
| [ ] | Content or scope of notified measure changed and text available from1: New deadline for comments (if applicable):  |
| [ ] | Interpretive guidance issued and text available from1:  |
| [ ] | Other:  |

**Description:** Ukraine notifies the adoption of the Resolution of the Cabinet of Ministers of Ukraine No. 1121 "Some Іssues of Safety and Verification of Medicinal Products" of 26 September 2024.

The Resolution was published and entered into force on 03 October 2024.

The Resolution also stipulates that Clauses 2, 5-8, 11, 12, 14-18 of the Regulation on the National System of Verification of Medicinal Products shall be applied by state authorities and business entities from 03 October 20024;

other clauses of the Regulation on the National System of Verification of Medicinal Products and the Procedure for Application of Safety Features to the Packaging of Medicinal Products, Determining Their Characteristics, Means of Verification and Setting Requirements for Encryption, Structure and Format of Information To Be Contained in Safety Features, are applied by business entities:

* on a voluntary basis from 01 January 2026, but not before the availability of the relevant technical capability in the national system of verification of medicinal products. The technical capability of the national system of verification of medicinal products will be effective from the date of publication on the website of the National Agency for Verification of Medicinal Products of the information on the commissioning of the centralised data warehouse of the national system of medicinal products verification;
* mandatory from 01 January 2028.

Medicinal products put into circulation before 01 January 2028 without the application of safety features on their packaging may be sold in Ukraine until their expiry date.

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1. This information can be provided by including a website address, a pdf attachment, or other information on where the text of the final/modified measure and/or interpretive guidance can be obtained. [↑](#footnote-ref-1)