NOTIFICATION

Addendum

The following communication, dated 23 September 2022, is being circulated at the request of the delegation of Brazil.

**\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Title:** Draft resolution number 730, 14 October 2019.

|  |  |
| --- | --- |
| **Reason for Addendum:** | |
| [ ] | Comment period changed - date: |
| [X] | Notified measure adopted - date: 1 March 2023 |
| [ ] | Notified measure published - date: |
| [ ] | Notified measure enters into force - date: |
| [ ] | Text of final measure available from[[1]](#footnote-1): |
| [ ] | Notified measure withdrawn or revoked - date:  Relevant symbol if measure re-notified: |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:** The Draft resolution number 730, 14 October 2019 - previously notified through G/TBT/N/BRA/928 - which proposes the updating of the Resolution – RDC number 185, 22 October 2001, which establishes the risk classification, market authorization and technical requirements for labelling and usage instructions of medical devices, was adopted as Resolution - RDC number 751, 15 September 2022

The final text is available only in Portuguese and can be downloaded at:

<http://antigo.anvisa.gov.br/documents/10181/5672055/RDC_751_2022_.pdf/37b2d641-82ec-4e64-bb07-4fc871936735>

**\_\_\_\_\_\_\_\_\_\_**

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)