NOTIFICATION

Addendum

The following communication, dated 17 November 2021, is being circulated at the request of the delegation of Canada.

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**Title:** Notice of Intent to Amend the Prescription Drug List (PDL): Brimonidine

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| **Reason for Addendum:** |
| [  ] | Comment period changed - date:  |
| [X] | Notified measure adopted - date: 2 November 2021 |
| [X] | Notified measure published - date: 15 November 2021 |
| [  ] | Notified measure enters into force - date:  |
| [X] | Text of final measure available from[[1]](#footnote-1): <https://www.canada.ca/en/health-canada/services/drugs-health-products/drug-products/prescription-drug-list/notices-changes/intent-amend-brimonidine.html> <https://www.canada.ca/fr/sante-canada/services/medicaments-produits-sante/medicaments/liste-drogues-ordonnance/avis-concernant-modifications/intention-modifier-brimonidine.html> |
| [  ] | Notified measure withdrawn or revoked - date: Relevant symbol if measure re-notified:  |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 purpose of this Notice of Intent to Amend is to notify that, as a result of a scientific review and public consultation, Health Canada will add a qualifier to brimonidine on the Prescription Drug List to enable sale of certain non-prescription products. Both the human and veterinary parts of the Prescription Drug List are to be revised. Health Canada has conducted a scientific review of brimonidine tartrate against a set of established and publicly available criteria outlined in section C.01.040.3 of the *Food and Drug Regulations* that led to this decision. The wording of the qualifier has not changed from that was proposed in the 1 September 2021 Notice of Consultation, notified under G/TBT/N/CAN/653. This revision will be in effect six months from the date that the Notice of Intent to Amend is posted on the Health Canada website.

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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)