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 தேசிய மருந்துகள் ஒழுங்குமுறை அதிகாரசபை
 National Medicines Regulatory Authority



සෞඛ්‍ය අමාත්‍යාංශය | சுகாதார அமைச்சு | Ministry of Health

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திகதி 04/03/2026

Notice on Labelling Requirement

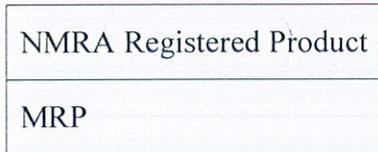
To All Marketing Authorization Holders of Medicines

Regulatory Strengthening of Controls on Falsified Products

The National Medicines Regulatory Authority has decided to implement certain changes to the labelling requirements for the medicines in order to enhance the protection against falsified products.

All commercial packs shall bear a printed or non-peelable sticker displaying the wording “NMRA Approved Product” together with the Maximum Retail Price (MRP). These details may appear either on separate stickers or on a single label, provided that the two statements are clearly distinguishable, preferably separated by a line.

A specimen is given below as an example:



The sticker shall be implemented immediately for all consignments released to the market with effect from 1st of April 2026. The batches already released to the Market are not affected by this decision.

We expect your fullest cooperation to implement these control measures and help prevention of falsified products getting into the market.

Thank you

Chief Executive Officer
National Medicines Regulatory Authority

Dr. Prabhath Werawatte
 Chief Executive Officer
 National Medicines Regulatory Authority
 No. 130, W.A.D. Ramanayake Mawatha,
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