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தேசிய மருந்துகள் ஒழுங்குமுறை அதிகாரசபை  
National Medicines Regulatory Authority



සෞඛ්‍ය අමාත්‍යාංශය | சுகாதார அமைச்சு | Ministry of Health

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Notice on Labelling Requirement-Amendment 01

**To All Marketing Authorization Holders of Medicines**

**Regulatory Strengthening of Controls on Falsified Products**

The National Medicines Regulatory Authority 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 Registered 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:

NMRA Registered Product
MRP

The sticker shall be implemented immediately for all consignments released to the market with effect from 1<sup>st</sup> 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,  
Colombo - 02

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රාජ්‍ය ඉංජිනේරු සංස්ථා ගොඩනැගිල්ල (2 වැනි මහල),  
අංක 130, ඩබ්ලිව්. ඒ. ඩී. රාමනායක මාවත,  
කොළඹ 02,  
ශ්‍රී ලංකා.

தேசிய மருந்துகள் ஒழுங்குமுறை ஆணையம்,  
இலங்கை அரசு பொறியியல் கழகக் கட்டிடம் (2வது மாடி),  
இல. 130, டபிள்யூ.ஏ.டி. ரமநாயக்க மாலத்தை,  
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