

Message 001

Communication from the Commission - TRIS/(2026) 0240

Directive (EU) 2015/1535

Notification: 2026/0027/ES

Notification of a draft text from a Member State

Notification – Notification – Notifizierung – Нотификация – Oznámení – Notifikation – Γνωστοποίηση – Notificación – Teavitamine – Ilmoitus – Obavijest – Bejelentés – Notifica – Pranešimas – Paziņojums – Notifika – Kennisgeving – Zawiadomienie – Notificação – Notificare – Oznámenie – Obvestilo – Anmälan – Fógra a thabhairt

Does not open the delays - N'ouvre pas de délai - Kein Fristbeginn - Не се предвижда период на прекъсване - Nezahtuje prodlení - Fristerne indledes ikke - Καμμία έναρξη προθεσμίας - No abre el plazo - Viivituste perioodi ei avata - Määräaika ei ala tästä - Ne otvara razdoblje kašnjenja - Nem nyitja meg a késésekét - Non fa decorrere la mora - Atidējimai nepradedami - Atlikšanas laikposms nesākas - Ma jiftaħ il-perijodi ta' dewmien - Geen termijnbegin - Nie otwiera opóźnień - Não inicia o prazo - Nu deschide perioadele de stagnare - Nezačína oneskorenia - Ne uvaja zamud - Inleder ingen frist - Ní osclaíonn sé na moilleanna

MSG: 20260240.EN

1. MSG 001 IND 2026 0027 ES EN 23-01-2026 ES NOTIF

2. Spain

3A. Ministerio de Asuntos Exteriores, Unión Europea y Cooperación

Dirección General de Coordinación del Mercado Interior y Otras Políticas Comunitarias

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3B. Área de Biocidas y Productos Químicos

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4. 2026/0027/ES - C00C - CHEMICALS

5. Draft Royal Decree regulating the registration and conditions for the authorisation, manufacture, marketing and use of biocidal products

6. The draft applies to biocidal products, which are regulated by Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012. Specifically, it updates current Spanish legislation in a way that complements the aforementioned Regulation.

7.

8. It updates the legal framework governing the registration of biocidal products in Spain.

It updates the specific conditions for the manufacture, packaging, storage, marketing and use of biocidal products within national territory.

It updates the training requirements for carrying out treatments with biocidal products in accordance with the new national educational requirements.

It updates the sanctions regime in accordance with the Biocidal Products Regulation.

9. Spanish legislation on biocidal products is currently scattered across several legislative acts and is outdated in some of the aspects regulated by Regulation (EU) No 528/2012. These include aspects relating to the official register of biocidal products, the official register of biocidal establishments and services, the official logbook of biocidal product movements, as well as the training of applicators that must be updated to reflect the new educational requirements regarding training.

Furthermore, the general conditions for the manufacture, storage, marketing and use of biocidal products as established in Royal Decree 3349/1983 of 30 November, must be updated in accordance with current practice.

In addition, the draft updates the chapter on infringements and sanctions in accordance with the obligations resulting from the Biocidal Products Regulation.

10. References to the basic texts:

11. No

12.

13. No

14. No

15. Yes

16.

TBT aspects: No

SPS aspects: No

European Commission

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