Message 001

Communication from the Commission - TRIS/(2026) 2621

Directive (EU) 2015/1535

Notification: 2026/0512/AT

Notification of a draft text from a Member State

Notification – Notification – Notifzierung – 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 - Nezahajuje 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 késéseket - Non fa decorrere la mora - Atidėjimai nepradedami - Atlikšanas laikposms nesākas - Ma jiftaħx 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: 20262621.EN

1. MSG 001 IND 2026 0512 AT EN 18-09-2026 AT NOTIF

2. Austria

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4. 2026/0512/AT - SERV - INFORMATION SOCIETY SERVICES

5. Regulation of the Federal Minister for Labour, Social Affairs, Health, Care and Consumer Protection amending the 2015 ELGA Regulation (Amendment to the ELGA Implementation Guidelines 2026)

6. Electronic health services/Defining the content, structure, format and coding of ELGA health data in implementation guidelines

7.

8.
The purpose of this draft is to amend the 2015 ELGA Regulation. The aim of the project is the operationalisation of a technical standard (CDA) for electronic medical records (ELGA). It mainly covers the following measures:
- Regulation of updated version of a previously specified implementation guideline
- Regulation of a new implementation guideline

9. Documents and data in ELGA must comply with the international CDA (Clinical Document Architecture) standard of the standardisation organisation HL7 in terms of technology and content, according to the resolutions of the ELGA system partners (government, federal provinces and social insurance). However, standards cannot be directly implemented, but need to be put into practice in a number of ways. This is done by means of so-called implementation guidelines. The new implementation guideline creates the prerequisites for ELGA-compliant preparation and use of health advice reports from the 1450 hotline. The existing ‘Care Report Implementation Guideline’ is to be updated.

9a. The prescribed implementation guidelines set out technical standards for the purposes of creating ELGA documents with a view to ensuring interoperability. These standards comply with the international CDA (Clinical Document Architecture) standard established by the HL7 standards body in terms of both technical specifications and content.

9b. Technical interoperability could not be ensured without standard specification. All health service providers that process ELGA documents must comply with these uniform standards.

9c. The requirement of uniform, international, technical standards is an appropriate measure to ensure the interoperability of electronic medical records (ELGA).

10. Reference to the basic texts: 2023/0439/AT

Basic texts have been forwarded under a previous notification:
2023/0439/AT

11. No

12.

13. No

14. No

15. Yes

16.
TBT aspects: No

SPS aspects: No

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