**ORDINARY LEGISLATIVE procedure**

**(First reading without prior interinstitutional negotiations)**

**Follow up to the European Parliament legislative resolution on the proposal for a regulation of the European Parliament and of the Council laying down Union procedures for the authorisation and supervision of medicinal products for human use and establishing rules governing the European Medicines Agency, amending Regulation (EC) No 1394/2007 and Regulation (EU) No 536/2014 and repealing Regulation (EC) No 726/2004, Regulation (EC) No 141/2000 and Regulation (EC) No 1901/2006**

**1. Rapporteur:** Tiemo WÖLKEN (S&D / DE)

**2. Reference numbers:** 2023/0131 (COD) / A9-0141/2024 / P9\_TA(2024)0221

**3. Date of adoption of the resolution:** 10 April 2024

**4. Legal basis:** Article 114(1) and Article 168(4), point (c) of the Treaty on the Functioning of the European Union

**5. Competent Parliamentary Committee:** Committee on Environment, Public Health and Food Safety (ENVI)

**6. Commission's position:** takes note of the amendments proposed by the European Parliament, while reserving its detailed position on these until the opening of interinstitutional negotiations.