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Summary of European Union decisions on marketing authorisations in respect of medicinal products from 1 January 2026 to 31 August 2026

(Published pursuant to Article 13 or Article 38 of Regulation (EC) No 726/2004 of the European Parliament and of the Council ([1]) or Article 5 of Regulation (EU) 2019/6 of the European Parliament and of the Council ([2]))

(C/2026/5454)

— **Issuing of a marketing authorisation** (Article 13 of Regulation (EC) No 726/2004 of the European Parliament and of the Council): **Accepted**

Date of the decision	Name of the medicinal product	INN (International Non-Proprietary Name)	Holder of the marketing authorisation	Number of the entry in the Community Register	Pharmaceutical form	ATC code (Anatomical Therapeutic Chemical Code)	Date of notification
20.8.2026	Nylaspeg	pegfilgrastim	QILU PHARMA SPAIN S.L. Paseo de la Castellana 40, planta 8, 28046 Madrid, España	EU/1/26/2051	Solution for injection	L03AA13	11.9.2026
28.8.2026	Denosumab Ascend	denosumab	Ascend GmbH Sebastian-Kneipp-Strasse 41, Niederursel, 60439 Frankfurt Am Main, Hesse, Deutschland	EU/1/26/2052	Solution for injection	M05BX04	18.9.2026

([1]) OJ L 136, 30.4.2004, p. 1, ELI: https://data.europa.eu/eli/reg/2004/726/oj.
([2]) OJ L 4, 7.1.2019, p. 43, ELI: https://data.europa.eu/eli/reg/2019/6/oj.

— **Modification of a marketing authorisation** (Article 13 of Regulation (EC) No 726/2004 of the European Parliament and of the Council): **Accepted**

Date of the decision	Name of the medicinal product	Holder of the marketing authorisation	Number of the entry in the Community Register	Date of notification
26.5.2026	Artesunate Amivas	Amivas Ireland Ltd Suite 5, Second Floor, Station House, Railway Square, Waterford, Ireland	EU/1/21/1582	8.6.2026
26.5.2026	Foclivia	Seqirus S.r.l. Via del Pozzo 3/A, S. Martino, 53035 Monteriggioni (SI), Italia	EU/1/09/577	28.5.2026
26.5.2026	Lacosamide UCB	UCB Pharma S.A. Allée de la Recherche 60, 1070 Bruxelles, Belgique/Researchdreef 60, 1070 Brussel, België	EU/1/19/1383	1.6.2026
26.5.2026	Prevenar 20	Pfizer Europe MA EEIG Boulevard de la Plaine 17, 1050 Bruxelles, Belgique/ Pleinlaan 17, 1050 Brussel, België	EU/1/21/1612	27.5.2026
26.5.2026	ZOLSKETIL pegylated liposomal	Accord Healthcare S.L.U. World Trade Center, Moll de Barcelona, s/n, Edifici Est, 6ª planta, 08039 Barcelona, España	EU/1/22/1629	27.5.2026
17.8.2026	Buprenorphine Neuraxpharm	Neuraxpharm Pharmaceuticals, S.L. Avda. Barcelona 69, 08970 Sant Joan Despí - Barcelona, España	EU/1/24/1809	26.8.2026
20.8.2026	Sitagliptin SUN	Sun Pharmaceutical Industries Europe BV Polarisavenue 87, 2132 JH Hoofddorp, Nederland	EU/1/21/1598	5.10.2026

— **Modification of a marketing authorisation** (Article 38 of Regulation (EC) No 726/2004 of the European Parliament and of the Council; Article 5 of Regulation (EU) 2019/6 of the European Parliament and of the Council): **Accepted**

Date of the decision	Name of the medicinal product	Holder of the marketing authorisation	Number of the entry in the Community Register	Date of notification
5.6.2026	STARTVAC	Laboratorios Hipra, S.A. Avda. La Selva, 135, 17170 Amer (Girona), España	EU/2/08/092	12.6.2026

Anyone wishing to consult the public assessment report on the medicinal products in question and the decisions relating thereto is invited to contact:

European Medicines Agency
Domenico Scarlattilaan 6
1083 HS Amsterdam
NETHERLANDS