Summary of European Commission Decisions on authorisations for the placing on the market for the use and/or for use of substances listed in Annex XIV to Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH)

(Published pursuant to Article 64(9) of Regulation (EC) No 1907/2006 ([1]))

(Text with EEA relevance)

(2023/C 78/06)

Decision granting an authorisation

Reference of the decision ([1])	Date of decision	Substance name	Holder of the authorisation	Authorisation number	Authorised use	Date of expiry of review period	Reasons for the decision
C(2023) 1180	23 February 2023	4-(1,1,3,3-tetramethylbutyl)phenol, ethoxylated ('4-tert-OPnEO') EC No.-; CAS No.-	Wallac Oy, Mustionkatu 6, 20750 Turku, Finland	REACH/23/5/0	Formulation of 4-tert-OPnEO (as Triton X-100) for use in the assay buffer for the GSP® Neonatal GALT kit used for the semi-quantitative determination of galactose-1-phosphate uridyl transferase (GALT) activity	4 January 2026	In accordance with Article 60(4) of Regulation (EC) No 1907/2006, the socio-economic benefits outweigh the risk to human health and the environment from the use of the substance and there are no suitable alternative substances or technologies.
				REACH/23/5/1	In the assay buffer of the GSP® Neonatal GALT kit used for the semi-quantitative determination of galactose-1-phosphate uridyl transferase (GALT) activity		

([1]) The decision is available on the European Commission website at: Authorisation (europa.eu).

([1]) OJ L 396, 30.12.2006, p. 1.