**Official request form for addition/amendments to the EU reference dates (EURD) list**

**This request form should be sent to** [EURDlist@ema.europa.eu](mailto:EURDlist@ema.europa.eu) **only.**

**REQUEST DETAILS**

**Type of request:** Please select request type

**EURD reference date:** Click here to enter a date.

**Name of active substance or combination of active substances:** *Please only enter 1 request per form*

Click here to enter text.

**Grounds for request:**

Click here to enter text.

**Please provide the list of relevant authorised medicinal products from Article 57 database, stating the product names and EV codes in the email. If more than 10 EV codes are concerned, please provide the product names and EV Codes in an Excel spreadsheet, one EV Code per cell and attach it in the email along with the form.**

Please note:

* If relevant PSUSA is starting or about to start, the request can be made in the context of PSUSA assessment (e.g. comments to the PAR)
* This document must be attached as it is to the email. Do not scan it. No signature is required on the document
* Please note that each EURD list amendment is only valid 6 months after its publication. Please take this into consideration for understanding/planning timelines for amendments
* Requests for additions/amendments of the EURD list require the agreement of the Member States in which each active substance/combination is authorised, the advice of the Granularity and Periodicity Advisory Group (GPAG) to PRAC and the PRAC Recommendation. Moreover, the CHMP and CMDh need to adopt any changes to the EURD list. A long timeframe is therefore required, for each request to be processed and accepted/rejected