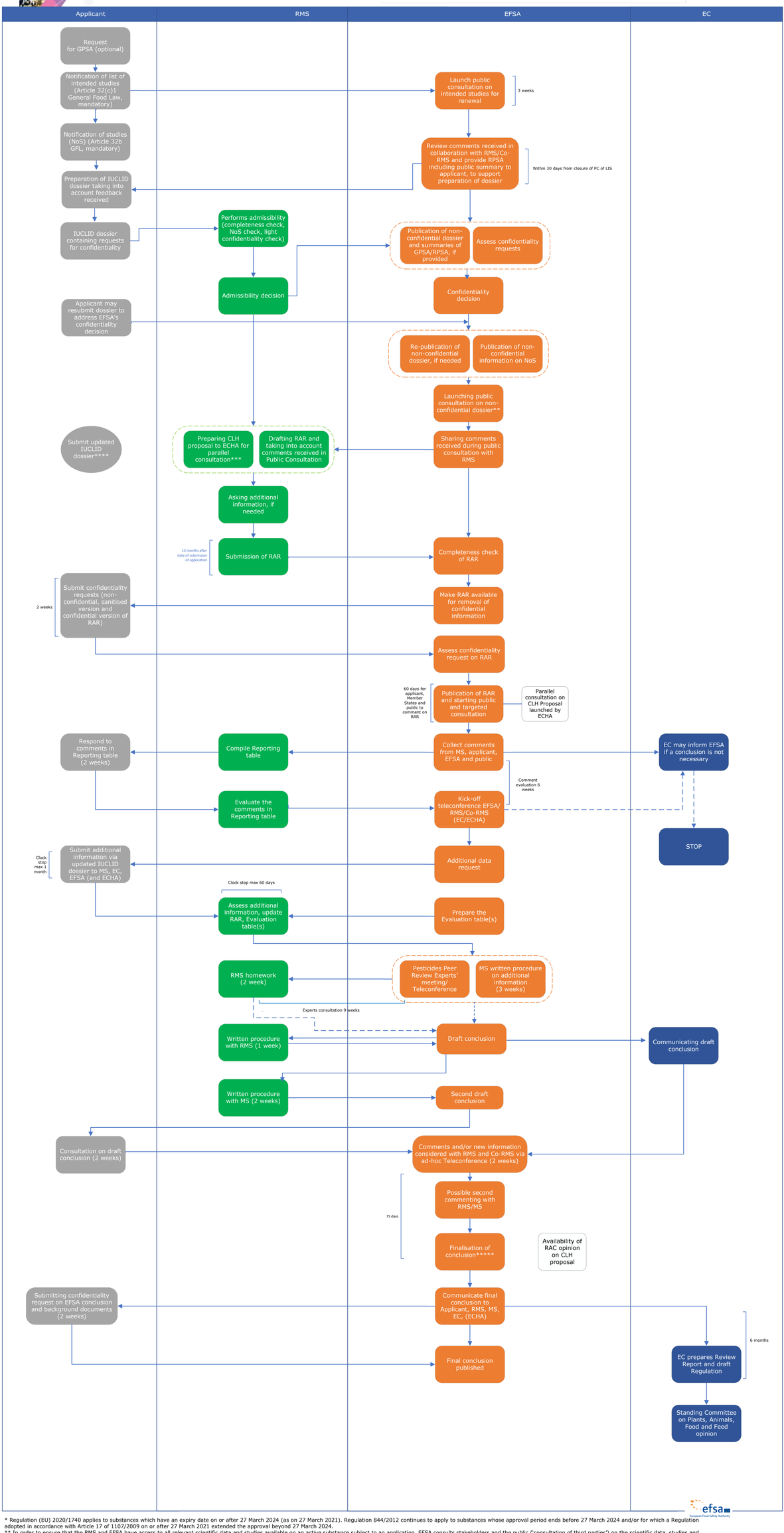
Renewal of approval of pesticide active substances under Regulation (EU)2020/1740*



* Regulation (EU) 2020/1740 applies to substances which have an expiry date on or after 27 March 2024 (as on 27 March 2021). Regulation 844/2012 continues to apply to substances whose approval period ends before 27 March 2024 and/or for which a Regulation adopted in accordance with Article 17 of 1107/2009 on or after 27 March 2021 extended the approval beyond 27 March 2024.

** In order to ensure that the RMS and EFSA have access to all relevant scientific data and studies available on an active substance subject to an application, EFSA consults stakeholders and the public ('consultation of third parties') on the scientific data, studies and other information part of, or supporting, the submitted application to identify whether other relevant scientific data or studies are available.

***To comply with the Regulation (EU) No 2020/1740, at the latest at the time of submission of the RAR, the RMS shall submit a proposal to ECHA to obtain an opinion on a harmonised classification of the active substance at least for the hazard classes defined in Article 11(9) of Commission Implementing Regulation (EU) 2020/1740, or to confirm the existing classification, where applicable, or for re-classification of the active substance in accordance with the criteria of Regulation (EC) No 1272/2008. This is to facilitate that the ECHA Risk Assessment Committee (RAC) Opinion could be available in time for consideration in the EFSA peer review.

**** Applicants are requested to update their dossiers during the admissibility check of the application as well as during the risk assessment by RMS and during EFSA peer-review. However, to the extent possible these requests should be consolidated in order to trigger the smallest number of dossier updates possible.

***** Overall timing to draft EFSA conclusion is within 5 months from the end of commenting + clock stop period + 2 weeks additional consultation + 75 days for finalisation or within two weeks from the adoption of the CLH opinion, whichever occurs later.

Co-RMS – Co-Rapporteur Member State
EC – European Commission
ECHA – European Chemicals Agency
GPSA – General Pre-submission Advice
LIS – List of Intended Studies
MS – Member State
NoS – Notification of Studies
PC – Public Consultation
RAR – (draft) Renewal Assessment Report
RMS – Rapporteur Member State
RPSA – Renewal Pre-submission Advice