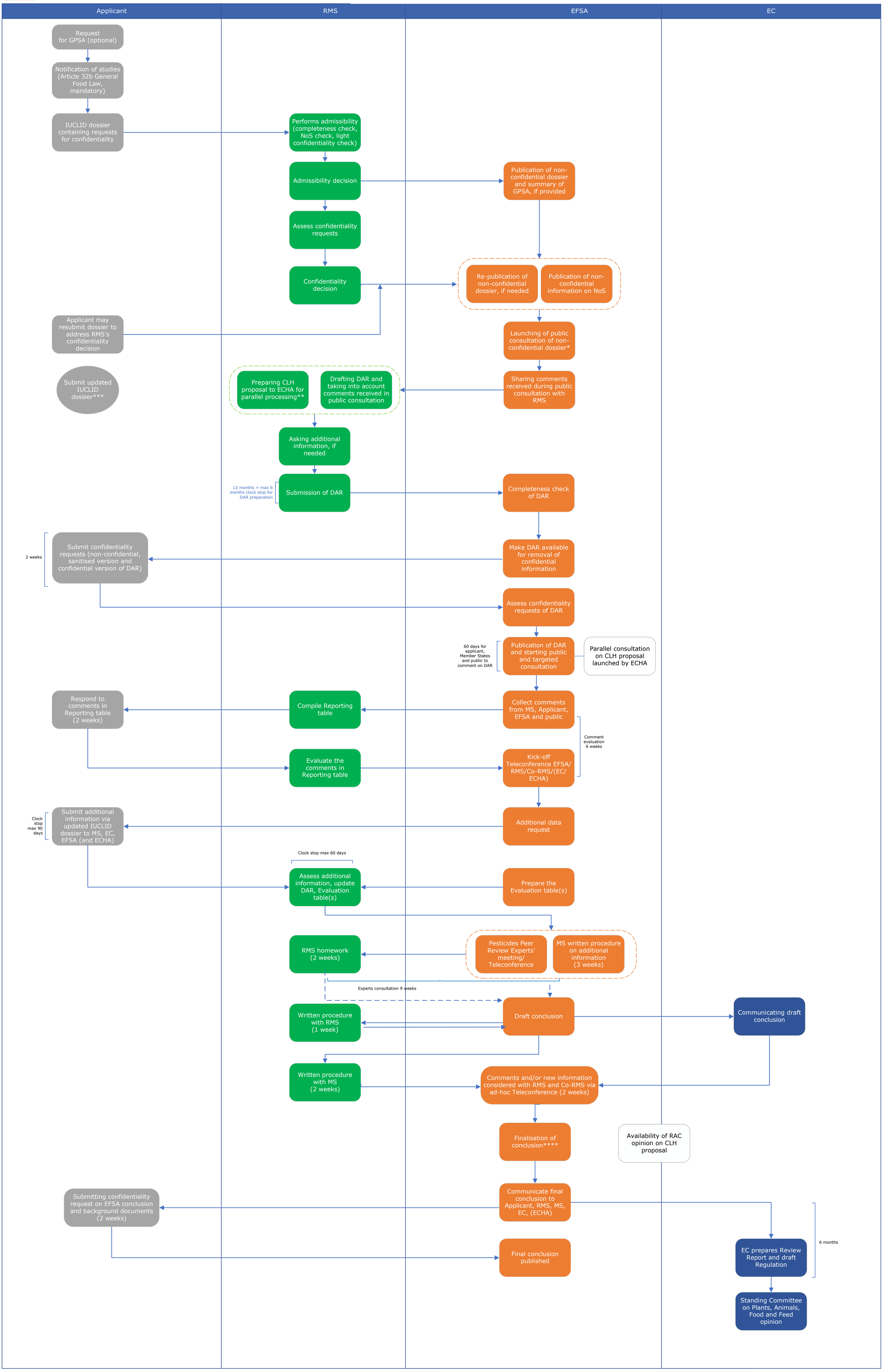
Approval of pesticide new active substances and amendment of approval conditions under Regulation 1107/2009 and (EU)2021/428



*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.
**The classification of pesticides is covered by Regulation (EC) No 1272/2008. Proposals for Harmonised Classification and Labelling (CLH) are handled at Risk Assessment Committee (RAC) of ECHA. To allow alignment of the EFSA peer review and ECHA classification processes, the RMS is strongly encouraged to submit a joint DAR/CLH report in parallel to both Agencies. EFSA and ECHA proceed to align procedures to facilitate that a RAC opinion on the harmonised classification of the active substance 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 finalise EFSA conclusion is within 5 months from the end of commenting + clock stop period (max 5 months).
DAR – Draft Assessment Report
EC – European Commission
ECHA – European Chemicals Agency
GPSA – General Pre-submission Advice
MS – Member State
NoS – Notification of Studies
RMS – Rapporteur Member State