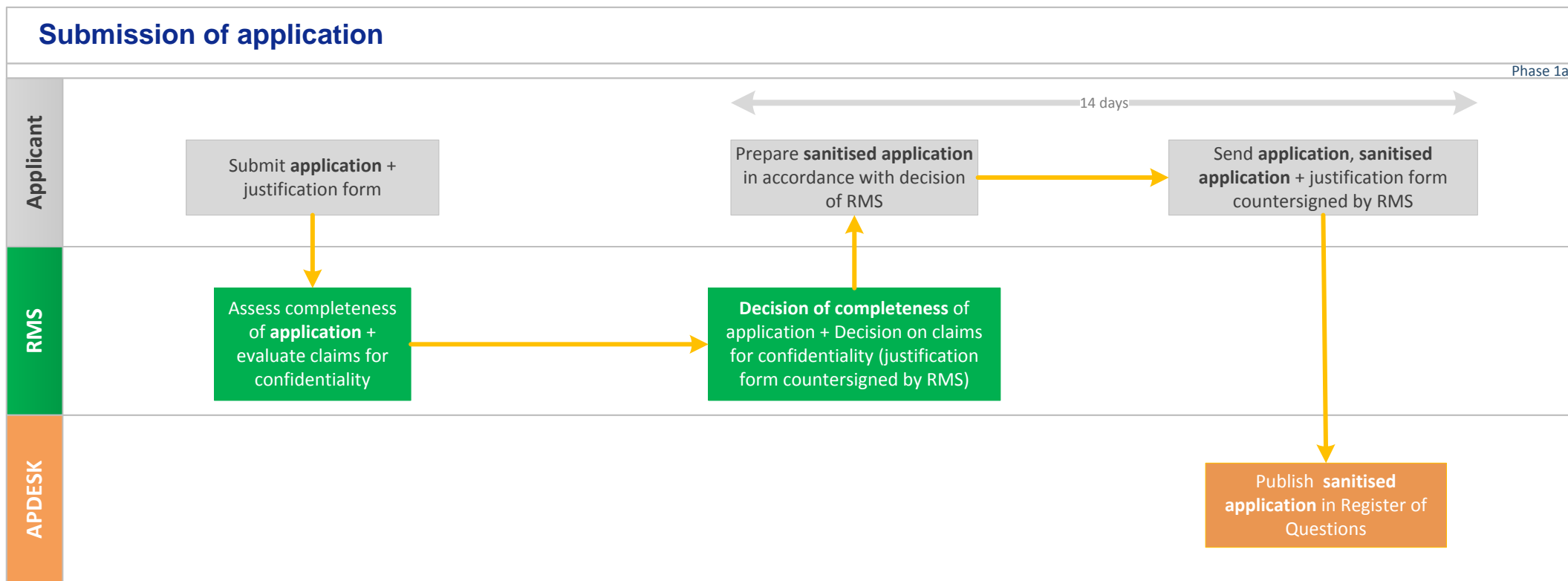




Applications helpdesk – Renewal of approval of active substances under Regulation EU 844/2012



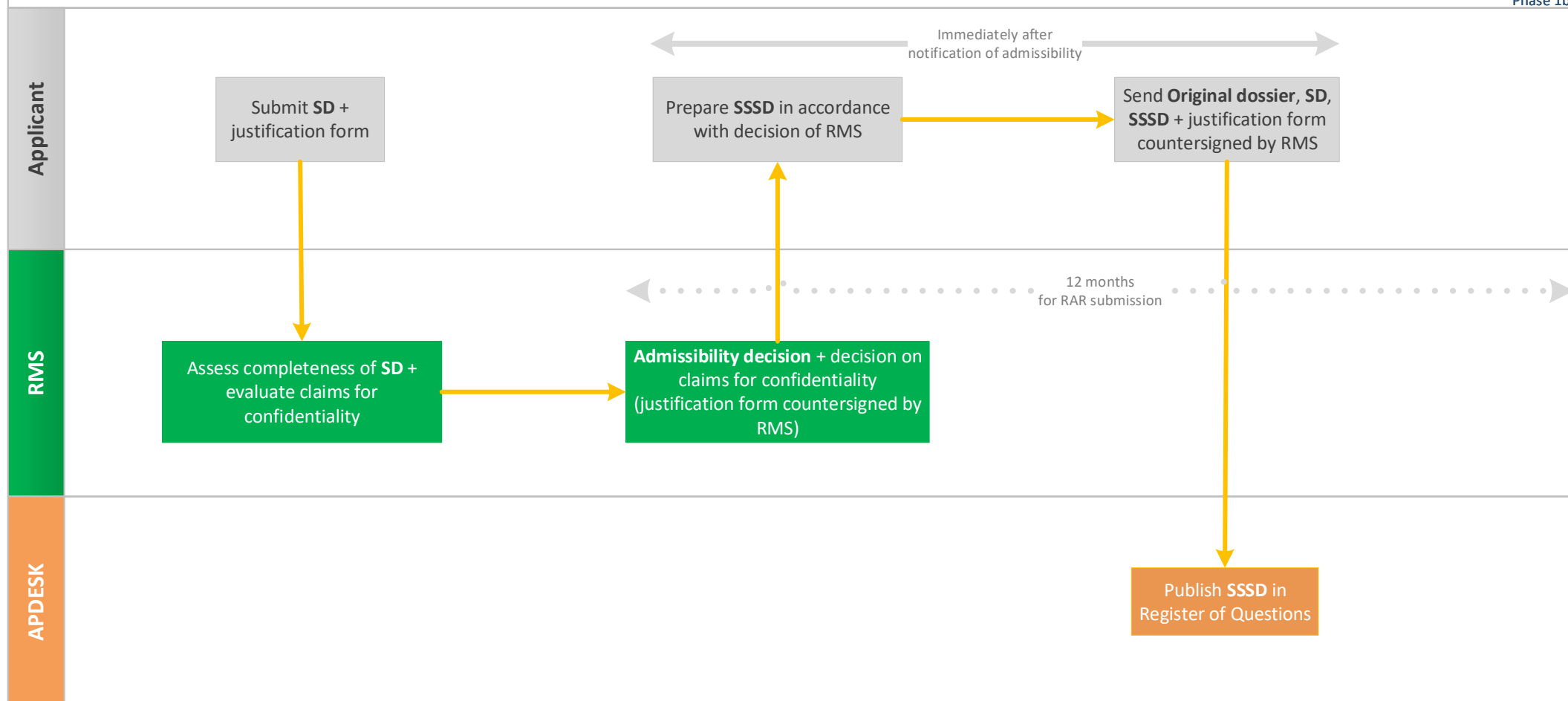
RMS: Rapporteur Member State
 EC: European Commission
 APDESK: Applications Desk Unit (EFSA)



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Submission of supplementary dossier

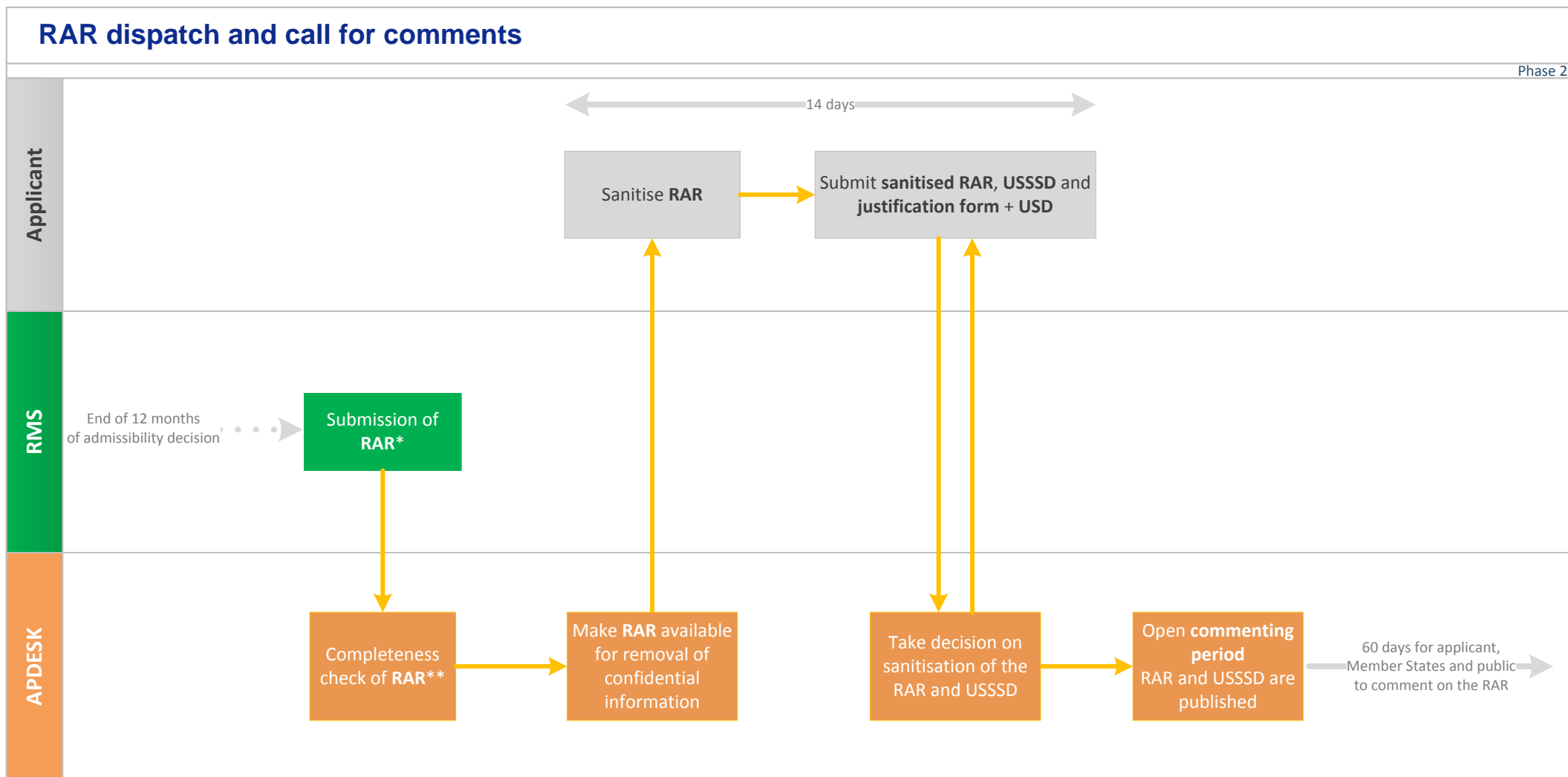
Phase 1b



SD: Supplementary Dossier
 SSSD: Sanitised Supplementary Summary Dossier
 RAR: Renewal Assessment Report



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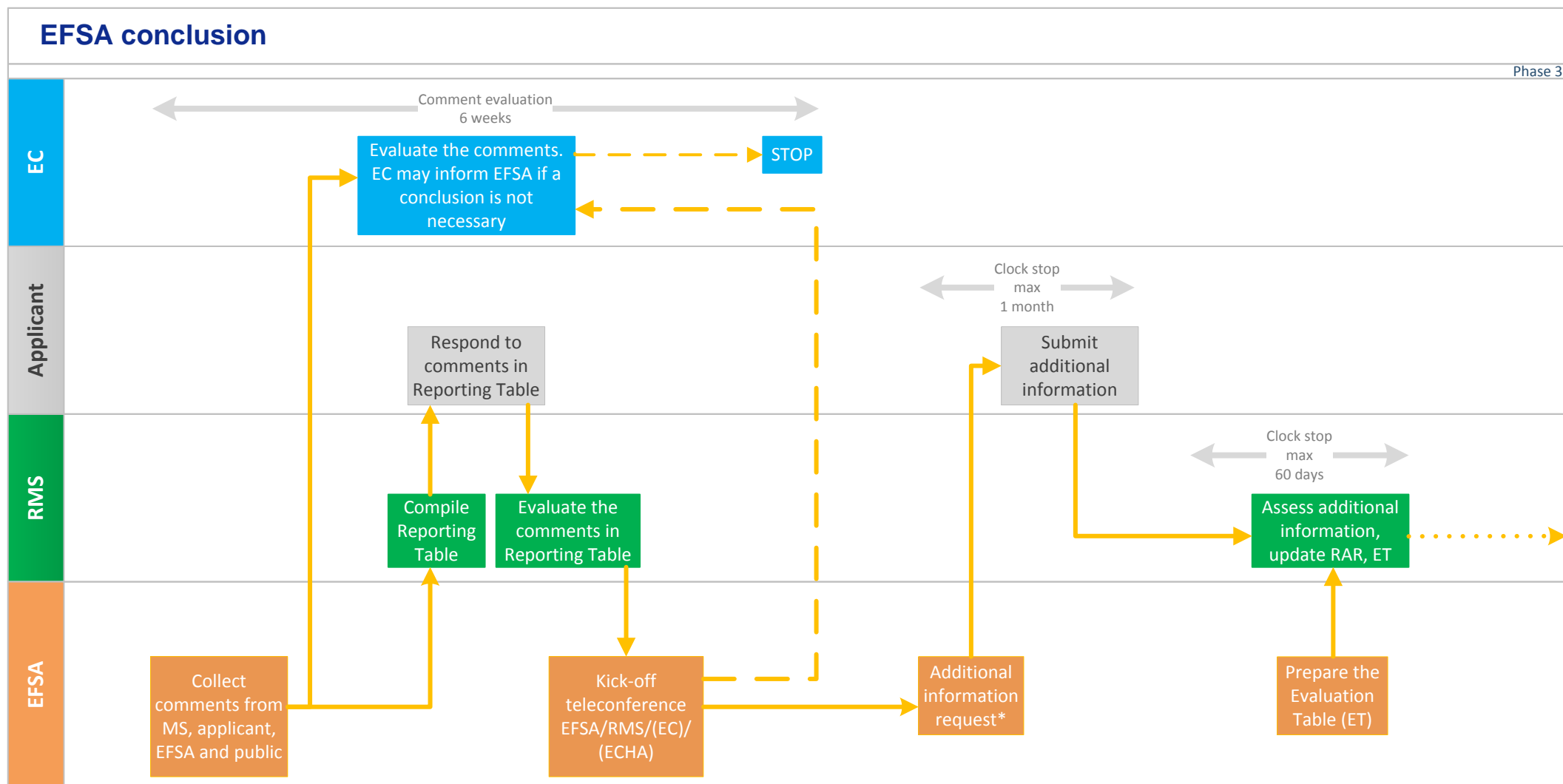
*After 10/11/2018 RAR should include updated ED assessment in line with new criteria.

**EFSA verifies that the RAR includes the ED assessment in line with the ECHA-EFSA Guidance. If this is missing, EFSA will ask for completion before proceeding with the peer-review.

USSSD: Updated Sanitised Supplementary Summary Dossier
 USD: Updated Supplementary Dossier



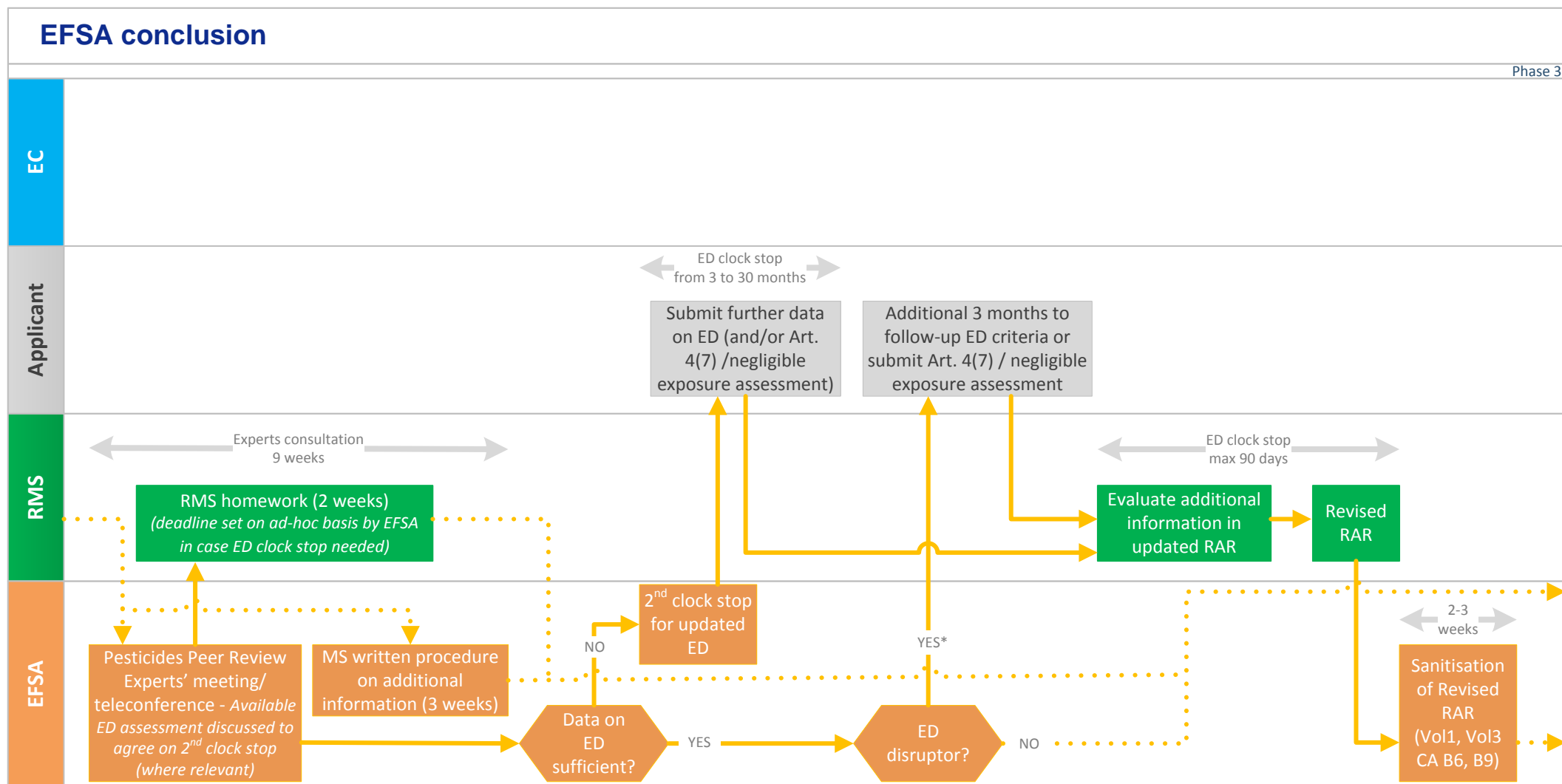
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* In case the initial RAR was submitted before 10/11/2018 and does not contain updated ED assessment, the clock stop of 1 month is used for giving opportunity to applicant to update the ED assessment in line with the ECHA-EFSA Guidance.



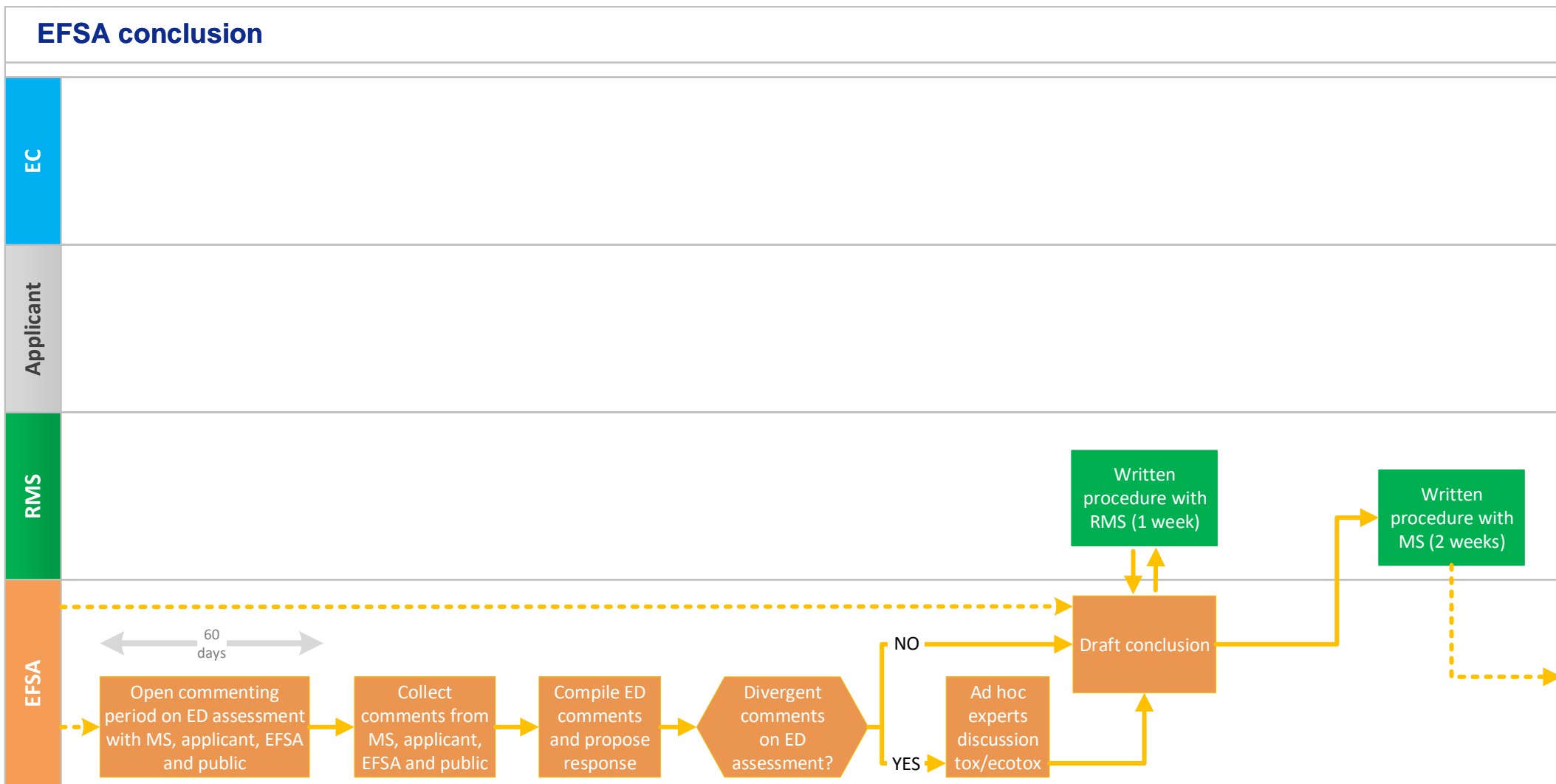
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*Criteria 3.6.5 and/or criteria 3.8.2 are met.



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