EUROPEAN COMMISSION
HEALTH AND CONSUMERS DIRECTORATE-GENERAL

Directorate E: Safety of Food chain
Unit E3: Chemicals, contaminants, pesticides

Reg. (EC) No 1107/2009
LIST OF IMPLEMENTING MEASURES

References Reg. 1107/2009	Deadlines	Measures
Art 84	14 June 2011	• Regulation containing the **list of the active substances already approved** at the moment of the adoption of that Regulation. • Regulation on **data requirements** for active substances {Art 8 (1)(b)} • Regulation on **data requirements** for PPP {Art 8 (1)(c)} • Regulation on **uniform principles** for risk assessment for PPP {Art 36} • Regulation containing requirements of the **labelling** of PPP {Art 6(1)}
Art 51 6a	14 December 2011	Report to European Parliament and Council on the establishment of a European **fund for minor uses**, accompanied, if appropriate, by a legislative proposal.
Art 67 1	14 December 2012	Report to European Parliament and Council on the costs and benefits of the traceability of the information concerning the PPP applications on agricultural products (if necessary, with legislative proposals).
Art 80 7	14 December 2013	List of **Candidates for substitution** included in Annex I of Directive 91/414/EEC (criteria point 4 Annex II)
Annex 2 3.6.5	14 December 2013	Draft measures concerning **specific scientific criteria for the determination of endocrine disrupting properties**.
Art 26	14 December 2014	Regulation establishing a **work programme** for **gradual review of synergists and safeners** on the market +relevant data requirements.
Art 82	14 December 2014	**Review clause** Report to European Parliament and Council on the application by MS of the provisions referred to in Art 36(3) and art 50(2), the division of the community in three zones and on the criteria for the approval substances, safeners and synergists as set in Annex II. Impact on the diversification and competitiveness of agriculture, on human health and environment (if necessary, with legislative proposals).
Art 30 3	14 June 2016	Deadline for application of **provisions on provisional authorisations**. Extensions possible by Comitology.
Art 62 3a	14 December 2016	Report + legislative proposal to European Parliament and Council on the effects of the provisions concerning **data protection** of tests and studies involving vertebrate animals. (if necessary, with legislative proposals)

References Reg. 1107/2009	Deadlines	Measures
Art 8 3	Before Application	**Format of the dossier** and Summary dossier to be submitted by industry. Advisory procedure.
Art 11 4	Before Application	**Format of the draft assessment report** of the rapporteur Member States on an active substance and of the assessment report on a plant protection product. Advisory procedure.
Art 13 4	Before Application & Regular up-date	Maintain a **list of approved active substances** electronically available to the public.
Art 22 3	N.D.	New criteria for approving an active substance as low-risk active substance.
Art 27 5 Art 81 2	N.D. but in line with derogation for MS in art. 81(2)	Detailed rules for the implementation of the provisions for **co-formulants**.
Art 21 1 Art 18	N.D.	**Renewal of approval**. **Work programme for renewal of approvals.** (Not Compulsory)
Art 58	N.D.	Detailed rules for the national authorization of **adjuvants** (data requirements, notification, evaluation, assessment and decision making criteria).
Art 68	N.D.	Regulation setting out provisions for **controls** carried out by the Member States.
Art 75 5	N.D.	The Commission shall publish and keep updated on its website a **list of the competent authorities** of the Member States
Art 77	N.D.	**Guidance documents** for the implementation of the Regulation such as explanatory notes or guidance documents on the content of the application concerning micro-organisms, pheromones and biological products. (Not Compulsory) Advisory procedure.
Art 29 4	N.D.	Harmonise **methods for analysis** of active substances, safeners and synergists and, where appropriate, any toxicologically, ecotoxicologically or environmentally relevant impurities and co-formulants (Not Compulsory)
Art 38 5	N.D.	Detailed rules on the assessment of **equivalence** of different sources of active substances. (Not Compulsory)
Art 39 4	N.D.	Detailed rules on **reporting and exchange of information** on applications for authorisation. (Not Compulsory)
Art 43 4	N.D.	Guidelines on **compliance checks** for plant plant protection products after renewal of approval of a substance. (Not Compulsory)
Art 50	N.D.	Guidelines on **comparative assessment** of plant protection products containing candidates for substitution (Not Compulsory)

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Art 52	N.D.	Further details and specific requirements for **parallel trade** in case of application where an authorisation has already been granted or for products for personal use only.
Art 54 5	N.D.	Detailed rules on **research and development**, in particular the maximum quantities of plant protection products that may be released during experiments or tests and the minimum data to be submitted. (Not Compulsory)
Art 39 4	N.D.	Detailed rules for reporting and exchange of information on applications for authorisation to facilitate **work-sharing.** (Not Compulsory)

N.D. = No dead lines in the Regulation