

Regulation (EC) No 1107/2009 of the European Parliament and of the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC

CHAPTER II

**ACTIVE SUBSTANCES, SAFENERS, SYNERGISTS AND CO-FORMULANTS**

*[<sup>F1</sup>SECTION 4*

**Registers**

*Article 27A*

**Approvals register**

- 1 The competent authorities must jointly establish and maintain a register of active substances, safeners, synergists, low-risk active substances, basic substances and candidates for substitution approved in accordance with this Regulation.
- 2 The entry on the register for each substance must contain the following information—
  - a the common name and identification numbers of the substance;
  - b the IUPAC name of the substance, where available;
  - c the minimum purity of the substance;
  - d in respect of each constituent territory to which the entry relates—
    - i) whether the substance has been approved as an active substance, safener, synergist, low-risk active substance, basic substance or candidate for substitution;
    - ii) the date of the approval decision;
    - iii) except in relation to approved basic substances, the expiration date of approval;
    - iv) information on any specific provisions, conditions or requirements in respect of the approved substance.
- 3 The register must contain a search facility.
- 4 The competent authorities must jointly make the register available for inspection by the public on a website maintained by one or more of the competent authorities.

*Article 27B*

**Unacceptable co-formulants register**

- 1 The competent authorities must jointly establish and maintain a register of co-formulants which are not acceptable for inclusion in a plant protection product in accordance with Article 27.
- 2 The entry on the register for each co-formulant must contain the following information—

---

*Status: Point in time view as at 31/12/2020.*

*Changes to legislation: There are currently no known outstanding effects for the Regulation (EC) No 1107/2009 of the European Parliament and of the Council, SECTION 4. (See end of Document for details)*

---

- a the common name of the co-formulant;
  - b the IUPAC name of the co-formulant (where available);
  - c the CAS number of the co-formulant (where available);
  - d the EC number of the co-formulant (where available);
  - e in respect of each constituent territory to which the entry relates—
    - i) the date of the decision that the co-formulant was not acceptable for inclusion in a plant protection product;
    - ii) the sunset date for the co-formulant;
    - iii) any conditions of restriction relating to the co-formulant;
    - iv) any other information regarding the co-formulant that the competent authority considers relevant.
- 3 The register must contain a search facility.
- 4 The competent authorities must jointly make the register available for inspection by the public on a website maintained by one or more of the competent authorities.]

---

**Textual Amendments**

- F1** Ch. 2 Section 4 inserted (31.12.2020) by [The Plant Protection Products \(Miscellaneous Amendments\) \(EU Exit\) Regulations 2019 \(S.I. 2019/556\)](#), regs. 1(1), **4(35)** (with Sch. 1); 2020 c. 1, Sch. 5 para. 1(1)

**Status:**

Point in time view as at 31/12/2020.

**Changes to legislation:**

There are currently no known outstanding effects for the Regulation (EC) No 1107/2009 of the European Parliament and of the Council, SECTION 4.