Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products (Text with EEA relevance)

CHAPTER IX

CANCELLATION, REVIEW AND AMENDMENT OF AUTHORISATIONS

Article 47

Obligation for notification of unexpected or adverse effects

1 On becoming aware of information concerning the authorised biocidal product, or the active substance(s) it contains, that may affect the authorisation, the holder of an authorisation shall without delay notify the competent authority [F1].... In particular, the following shall be notified:

a new data or information on the adverse effects of the active substance or biocidal product for humans, in particular vulnerable groups, animals or the environment;

b any data indicating the potential of the active substance for the development of resistance;

c new data or information indicating that the biocidal product is not sufficiently effective.

2 The competent authority [F2]..., shall examine whether the authorisation needs to be amended or cancelled in accordance with Article 48.

[F3]3 . . . . . .

Textual Amendments

F1 Words in Art. 47(1) omitted (31.12.2020) by virtue of The Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/720), reg. 1(2), **Sch. 2 para. 94(a)**; 2020 c. 1, Sch. 5 para. 1(1)

F2 Words in Art. 47(2) omitted (31.12.2020) by virtue of The Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/720), reg. 1(2), **Sch. 2 para. 94(b)**; 2020 c. 1, Sch. 5 para. 1(1)

F3 Art. 47(3) omitted (31.12.2020) by virtue of The Chemicals (Health and Safety) and Genetically Modified Organisms (Contained Use) (Amendment etc.) (EU Exit) Regulations 2019 (S.I. 2019/720), reg. 1(2), **Sch. 2 para. 94(c)**; 2020 c. 1, Sch. 5 para. 1(1)