

This section presents the implementing regulations for Directive 98/8/EC and Decisions, organised by theme. These texts are published in the Official Journal of the European Union (OJEU).

Decisions and Regulations are legal acts of the European Union and are directly applicable in Member States in all their provisions.

• Regulations applicable since 1st September 2013

[Commission implementing Regulation \(EU\) No 2019/1825 of 8 August 2019](#) 

amending Implementing regulation (EU) No 528/2012 to include concentrated apple juice as an active substance in Annex I thereto

[Commission implementing Regulation \(EU\) No 2019/1824 of 8 August 2019](#) 

amending Implementing regulation (EU) No 528/2012 to include cheese as an active substance in Annex I thereto

[Commission implementing Regulation \(EU\) No 2019/1823 of 8 August 2019](#) 

amending Implementing regulation (EU) No 528/2012 to include D- fructose as an active substance in Annex I thereto

[Commission implementing Regulation \(EU\) No 2019/1822 of 8 August 2019](#) 

amending Implementing regulation (EU) No 528/2012 to include honey as an active substance in Annex I thereto

[Commission implementing Regulation \(EU\) No 2019/1821 of 8 August 2019](#) 

amending Implementing regulation (EU) No 528/2012 to include powdered egg as an active substance in Annex I thereto

[Commission implementing Regulation \(EU\) No 2019/1820 of 8 August 2019](#) 

amending Implementing regulation (EU) No 528/2012 to include *saccharomyces cerevisiae* as an active substance in Annex I thereto

[Commission implementing Regulation \(EU\) No 2019/1819 of 8 August 2019](#) 

amending Implementing regulation (EU) No 528/2012 to include vinegar as an active substance in Annex I thereto

[Commission implementing Regulation \(EU\) No 2016/1802 of 11 October 2016](#) 

amending Implementing regulation (EU) No 414/2013 specifying a procedure for the authorisation of same biocidal products in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission delegated Regulation \(EU\) No 492/2014 of 7 March 2014](#) 

supplementing Regulation (EU) No 528/2012 of the European Parliament and of the Council as regards the rules for the renewal of authorisations of biocidal products subject to mutual recognition

[Regulation \(EU\) No 334/2012 of the European Parliament and of the Council of 11 March 2014](#) 

amending Regulation (EU) No 528/2012 concerning the making available on the market and use of biocidal products, with regards to certain conditions for access to the market

[Commission implementing Regulation \(EU\) No 88/2014 of 31 January 2014](#) 

specifying a procedure for the amendment of Annex I to Regulation (EU) No 528/2012 of the European Parliament and of the Council concerning the making available on the market and use of biocidal products

[Commission implementing Regulation \(EU\) No 564/2013 of 18 June 2013](#) 

on the fees and charges payable to the European Chemicals Agency pursuant to Regulation (EU) No 528/2012 of the European Parliament and of the Council concerning the making available on the market and use of biocidal products

[Commission implementing Regulation \(EU\) No 414/2013 of 6 May 2013](#) 

specifying a procedure for the authorisation of same biocidal products in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission implementing Regulation \(EU\) No 354/2013 of 18 April 2013](#) 

on changes of biocidal products authorised in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

[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

• **Phases of the work programme for the placing on the market of biocidal products**

[Commission Delegated Regulation \(EU\) No 2019/227 of 28 November 2018](#) 

amending Delegated Regulation (EU) No 1062/2014 as regards certain active substances/product- type combinations for which the competent authority of the United Kingdom has been designated as the evaluating competent authority

[Commission Delegated Regulation \(EU\) No 2019/157 of 6 November 2018](#) 

amending Delegated Regulation (EU) No 1062/2014 on the work programme for the systematic examination of all existing active substances contained in biocidal products referred to in Regulation (EU) No 528/2012 of the European Parliament and of the Council concerning the making available on the market and use of biocidal products

[Commission Delegated Regulation \(EU\) No 2017/698 of 3 February 2017](#) 

amending Delegated Regulation (EU) No 1062/2014 on the work programme for the systematic examination of all existing active substances contained in biocidal products referred to in Regulation (EU) No 528/2012 of the European Parliament and of the Council concerning the making available on the market and use of biocidal products

Regulations & decisions

Written by DPR Biocides - Anses - Last Updated Friday, 23 October 2020 10:14

[Corrigendum to commission Delegated Regulation \(EU\) No 1062/2014 of 4 August 2014](#) 

on the work programme for the systematic examination of all existing active substances contained in biocidal products referred to in Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission Delegated Regulation \(EU\) No 1062/2014 of 4 August 2014](#) 

on the work programme for the systematic examination of all existing active substances contained in biocidal products referred to in Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission Delegated Regulation \(EU\) No 736/2013 of 17 May 2013](#) 

amending Regulation (EU) No 528/2012 of the European Parliament and of the Council as regards the duration of the work programme for examination of existing biocidal active substances

• Other provisions

[Commission implementing decision \(EU\) No 2020/1038 of 15 July 2020](#) 


postponing the expiry date of approval of creosote for use in biocidal products for product-type 8

[Commission implementing decision \(EU\) No 2020/1037 of 15 July 2020](#) 

postponing the expiry date of approval of acrolein for use in biocidal products for product-type 12

[Commission implementing decision \(EU\) 2019/1958 of 25 November 2019](#) 

on a derogation from mutual recognition of the authorisation of a biocidal product containing hydrogen cyanide by Poland in accordance with Article 37 of Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission implementing decision \(EU\) No 2019/1969 of 26 November 2019](#) 

postponing the expiry date of approval of IPBC for use in biocidal products for product-type 8

[Commission implementing decision \(EU\) No 2019/1951 of 25 November 2019](#) 

postponing the expiry date of approval of tebuconazole for use in biocidal products for product-type 8

[Commission implementing decision \(EU\) No 2019/1950 of 25 November 2019](#) 

postponing the expiry date of approval of K-HDO for use in biocidal products for product-type 8

[Commission implementing decision \(EU\) 2019/641 of 19 April 2019](#) 

on the terms and conditions of the authorisation of a biocidal product family containing 1R-trans phenothrin referred by Ireland in accordance with Article 36 of Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission implementing decision \(EU\) 2017/810 of 10 May 2017](#) 

on a derogation from mutual recognition of the authorisation of a biocidal product containing boric acid by France in accordance with Article 37 of Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission implementing decision \(EU\) 2016/1943 of 4 November 2016](#) 

pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on the use of paraffin oil for coating eggs to control the population size of nesting birds

[Commission implementing decision \(EU\) 2016/904 of 8 June 2016](#) 

pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on propan-2-ol containing products used for hand disinfection

[Commission implementing decision \(EU\) 2016/903 of 8 June 2016](#) 

pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on a horse rug impregnated with permethrin used for the purpose of controlling nuisance insects in the environment of the horse

[Commission implementing decision \(EU\) No 2016/135 of 29 January 2016](#) 

postponing the expiry date of approval of flocoumafen, brodifacoum and warfarin for use in biocidal products for product-type 14

[Commission implementing decision \(EU\) No 2015/1985 of 4 November 2015](#) 

pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on an anti-viral tissue impregnated with citric acid

[Commission implementing decision \(EU\) No 2015/1737 of 28 September 2015](#) 

postponing the expiry date of approval of bromadiolone, chlorophacinone and coumatetralyl for use in biocidal products for product-type 14

[Commission implementing decision \(EU\) No 2015/655 of 23 April 2015](#) 

pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on a polydimethylsiloxane-based formulation placed on the market to control mosquitoes

[Commission implementing decision \(EU\) No 2015/646 of 23 April 2015](#) 

pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on bacterial cultures intended to reduce organic solids and to be placed on the market for that purpose

[Commission implementing decision \(EU\) No 2015/411 of 11 March 2015](#) 


pursuant to Article 3(3) of Regulation (EU) No 528/2012 of the European Parliament and of the Council on cationic polymeric binders with quaternary ammonium compounds incorporated in paints and coatings

[Commission implementing decision 2014/397/EU of 25 June 2014](#) 

postponing the expiry date of approval of difethialone and difenacoum for use in biocidal products for product-type 14

Regulations & decisions

Written by DPR Biocides - Anses - Last Updated Friday, 23 October 2020 10:14

[Commission decision 2014/395/EU of 24 June 2014](#) 

concerning the placing on the market for essential use of biocidal products containing copper

[Commission Delegated Regulation \(EU\) No 837/2013 of 25 June 2013](#) 

amending Annex III to Regulation (EU) No 528/2012 of the European Parliament and of the Council as regards the information requirements for authorisation of biocidal products