



## Rejected substances


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This section provides a non-exhaustive list of the non-approval decisions that are applicable to all European countries. They are specific to an active substance/product type combination.


[Corrigendum to commission implementing Decision \(EU\) 2018/1622 of 29 October 2018](#)   
on the non-approval of certain biocidal active substances pursuant to Regulation (EU) No 528/2012 of the European Parliament and of the Council


[Commission implementing Decision \(EU\) 2018/1985 of 13 december 2018](#)   
not approving "Willaertia magma C2c Maky" as an active substance for use in biocidal products for product-type 11

[Commission implementing Decision \(EU\) 2018/1622 of 29 October 2018](#)   
on the non-approval of certain biocidal active substances pursuant to Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission implementing Decision \(EU\) 2018/1251 of 18 September 2018](#)   
not approving empenethrin as an existing active substance for use in biocidal products for product-type 18

[Commission implementing Decision \(EU\) 2018/622 of 20 April 2018](#)   
not approving chlorophene as an existing active substance for use in biocidal products for product-type 3

[Commission implementing Decision \(EU\) 2018/619 of 20 April 2018](#)   
not approving PHMB (1415; 4.7) as an existing active substance for use in biocidal products for product-type 1, 5 and 6

[Commission implementing Decision \(EU\) 2017/1282 of 14 July 2017](#)   
not approving 2-méthyl-1,2-benzisothiazol-3(2H)-one as an existing active substance for use in

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biocidal products for product-type 13

[Commission implementing Decision \(EU\) 2017/802 of 10 May 2017](#) 

not approving PHMB (1600; 1.8) as an existing active substance for use in biocidal products for product-type 5

[Commission implementing Decision \(EU\) 2016/1950 of 4 November 2016](#) 

on the non-approval of certain biocidal active substances pursuant to Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission implementing Decision \(EU\) 2016/110 of 27 January 2016](#) 

not approving triclosan as an active substance for use in biocidal products for product-type 1

[Commission implementing Decision \(EU\) 2016/109 of 27 January 2016](#) 

not to approve PHMB (1600; 1.8) as an active substance for use in biocidal products for product-types 1, 6 and 9

[Commission implementing Decision \(EU\) 2016/108 of 27 January 2016](#) 

not approving 2-Butanone, peroxide as an active substance for use in biocidal products for product-types 1 and 2

[Commission implementing Decision \(EU\) 2016/107 of 27 January 2016](#) 


not approving cybutryne as an active substance for use in biocidal products for product-type 21

[Commission implementing Decision \(EU\) 2015/1736 of 28 September 2015](#) 

not approving triflumuron as an active substance for use in biocidal products for product-type 18

[Commission implementing Decision 2014/227/EU of 24 April 2014](#) 

on the non-approval of certain biocidal active substances pursuant to Regulation (EU) No 528/2012 of the European Parliament and of the Council

[Commission Decision 2013/204/EU of 27 April 2013](#) 

concerning the non-inclusion of formaldehyde for product-type 20 in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2013/85/EU of 14 February 2013](#) 


concerning the non-inclusion of certain substances in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2012/728/EU of 23 November 2012](#) 

concerning the non-inclusion of bifenthrin for product type 18 in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2012/257/EU of 11 May 2012](#) 

concerning the non-inclusion of naled for product type 18 in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2012/254/EU of 10 May 2012](#) 

concerning the non-inclusion of dichlorvos for product type 18 in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2012/78/EU of 1 February 2012](#) 

concerning the non-inclusion of certain substances in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2012/77/EU of 1 February 2012](#) 


concerning the non-inclusion of flufenoxuron for product type 18 in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2011/391/EU of 1 July 2011](#) 

concerning the non-inclusion of certain substances in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2010/675/EU of 8 November 2010](#) 

concerning the non-inclusion of certain substances in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2010/72/EU of 8 February 2010](#) 

concerning the non-inclusion of certain substances in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2010/71/EU of 8 February 2010](#) 

concerning the non-inclusion of diazinon in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2009/324/EC of 14 April 2009](#) 

concerning the non-inclusion of certain substances in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2009/322/EC of 8 April 2009](#) 

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concerning the non-inclusion of certain substances in Annexes I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2008/809/EC of 14 October 2008](#) 

concerning the non-inclusion of certain substances in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2008/681/EC of 28 July 2008](#) 

concerning the non-inclusion of certain substances in Annexes I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2007/597/EC of 27 August 2007](#) 

concerning the non-inclusion of guazatine triacetate in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market

[Commission Decision 2007/565/EC of 14 August 2007](#) 

concerning the non-inclusion in Annex I, IA or IB to Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market of certain substances to be examined under the 10-year work programme referred to in Article 16(2) thereof