BPR Article 65(3) reporting

1. General information

1. General information
1.1. Member State For which Memeber State* are you reporting? ("In this survey "Member State" refers to EU Member States, loeland, Liechtenstein, Norway and Switzerland)
O Austria
O Belgium
O Bulgaria
O Croatia
○ Cyprus
O Czech Republic
○ Denmark
○ Estonia
○ Finland
France
○ Germany
○ Greece
○ Hungary
○ Iceland
O Ireland
O Italy
○ Latvia

LuxembourgMaltaNetherlands

O Liechtenstein
O Lithuania

- O Norway
- O Poland
- Portugal
- Romania
- Slovakia
- Slovenia
- Spain
- Sweden
- Switzerland
- United Kingdom

1.2. Contact details of the person responsible for reporting

	Name	Organisation	Email address	Telephone number(s)
		Ministry of Environment Ministry		
L		of Environment		

1.3. Information on the Competent Authority (CA)

1.3.1. Competent Authorities involved in the implementation of the BPR

How many CAs are responsible for the implementation of the BPR in your Member State? Please do not include enforcement authorities here, as they are specifically covered in section 4.

MTES (Ministry for the Ecological and Inclusive Transition) and ANSES (French Agency for Food, Environmental and Occupational He alth α Safety)

1.3.2. Details of the Competent Autorities involved in the BPR implementation

	BPR competent authority involved	Website
Advice to applicants/helpdesks on active substances	ANSES + Ministry for the Ecological and Inclusive Transition	https://www.helpdesk- biocides.fr/
Advice to applicants/helpdesks on biocidal products	ANSES + Ministry for the Ecological and Inclusive Transition	https://www.helpdesk- biocides.fr/

Advice to applicants/helpdesks on treated articles	ANSES + Ministry for the Ecological and Inclusive Transition	https://www.helpdesk- biocides.fr/
Assessment of active substances	ANSES	https://www.anses.fr/fr
Assessment and authorisation of biocidal products	ANSES	https://www.anses.fr/fr
Other (e.g. authority in charge of setting up the whole organisational	Ministry for the Ecological and	https://www.ecologique-
framework for the BPR implementation, of adopting national legislation)	Inclusive Transition	solidaire.gouv.fr/

1.3.3. Other bodies involved in the implementation of the BPR

	Authority/organisation involved	Website
Poison centre	Toxicovigilance and poison control centre	http://www.centres-antipoison.net/paris/
Animal poison	Toxicovigilance and poison control	http://www.vetagro-sup.fr/centre-national-dinformations-toxicologiques-veterinaires-cnitv/
centre	centre	www.centre-antipoison-animal.com
Other		

2. Relevant national measures and Member State specific measures

2.1. Transitional period (Art. 89 BPR)

Do you have specific national measures or legislation for making available on the market of biocidal products during the transitional period?

Yes

O No

Please specify below the national regulation(s) and/or requirement(s) during the transitional period or refer to the corresponding link of the relevant website with the requested information. If available in English please include the link to the English version.

Please also indicate whether such regulation(s) and/or requirement(s) changed during the reporting period.

In order to make available a biocidal product on the french market during the transitional period, the following requirements are applicable (these requirements are also applicable after the transitional period, except for the labelling):

- before the placing on the market, the product has to be declared on a national database called "SIMMBAD"

Article L522-2 (point I), and article article R522-18 to R522-20 of the Environmental Code: https://www.legifrance.gouv.fr/affich Code.do;jsessionid=904F85F50756DF99C64D484AE63EA5B9.tplgfr38s_2?idSectionTA=LEGISCTA000027723154&cidTexte=LEGITEXT000006074220&da teTexte=20200416

https://www.legifrance.gouv.fr/affichCodeArticle.do;jsessionid=904F85F50756DF99C64D484AE63EA5B9.tplgfr38s_2?idArticle=LEGIARTI000032815736scidTexte=LEGITEXT000006074220sdateTexte=20160701scategorieLien=idsoldAction=

- to provide to the organisations in charge of toxicovigilance the information concerning its detailed composition, on a specific database called "SYNAPSE"

Article L522-2 (point II) and article R522-21 of the Environmental Code :

https://www.legifrance.gouv.fr/affichCode.do;jsessionid=904F85F50756DF99C64D484AE63EA5B9.tplgfr38s_2?idSectionTA=LEGISCTA000027723154&cidTexte=LEGITEXT000006074220&dateTexte=20200416

https://www.legifrance.gouv.fr/affichCodeArticle.do;jsessionid=904F85F50756DF99C64D484AE63EA5B9.tplgfr38s_2?idArticle=LEGIARTI000
032815699scidTexte=LEGITEXT000006074220sdateTexte=20160701scategorieLien=idsoldAction=snbResultRech=

- to report each year the quantities placed on the market for every product

Article L522-3 and article 522-22 of the Environmental Code :

https://www.legifrance.gouv.fr/affichCode.do;jsessionid=904F85F50756DF99C64D484AE63EA5B9.tplgfr38s_2?idSectionTA=LEGISCTA000027723154&cidTexte=LEGITEXT000006074220&dateTexte=20200416

https://www.legifrance.gouv.fr/affichCodeArticle.do;jsessionid=904F85F50756DF99C64D484AE63EA5B9.tplgfr38s_2?idArticle=LEGIARTI000032815691&cidTexte=LEGITEXT000006074220&dateTexte=20160701&categorieLien=id&oldAction=&nbResultRech=

- to follow specific labelling requirements :

Article L522-8 of the Environmental Code and decree of 19 may 2004 on the control of the placing on the market of biocidal active substances and the authorisation of the placing on the market of biocidal products

 $\label{lem:https://www.legifrance.gouv.fr/affichCode.do?idSectionTA=LEGISCTA000027723129&cidTexte=LEGITEXT000006074220&dateTexte=20200416 \\ \text{https://www.legifrance.gouv.fr/affichTexte.do?cidTexte=JORFTEXT000000251018&categorieLien=id} \\ \\$

2.2. Applicable fees

Do you have specific national measures or legislation regarding fees for BPR procedures?

- Yes
- O No

Please specify below the national regulation(s) and indicate the corresponding link to the relevant website with the requested information. If available in English please include the link to the English version

Arrêté du 22 novembre 2017 fixant le montant de la rémunération due au titre de l'approbation et de l'autorisation de mise sur le marché des substances et produits biocides: https://www.helpdesk-biocides.fr/files/PDF/docume nts_utiles/informatifs/fr/20171122_redevances_france.pdf

2.2.1. Fee amounts

Please provide information on the applicable fees for the procedures listed in the table below

	Fee amount
Evaluation of an active substance for approval	For one biocidal product type : 200 000€ For each additional product type : + 100 000€ When the AS is a micro-organism and for one biocidal product type : 120 000€ For each additional product type : + 60 000€
Evaluation of an active substance for Annex I inclusion	None
Authorisation of a biocidal product (BP)	For one biocidal product type and one user type : 40 000€ For each additional user type : + 8000€ For each additional product type : + 8000€ When the product contain at least one AS candidate to substitution : + 10 000 €
Authorisation of a BP family	For one biocidal product type and one user type: 80 000€ For each additional user type: + 16 000€ For each additional product type: + 16 000€ When the product contain at least one AS candidate to substitution: + 20 000 € For each biocidal product in the family: + 800€/product
Mutual recognition of an authorisation of a BP	For one biocidal product type and one user type : 15 000€ For each additional user type : + 3000€ For each additional product type : + 3000€
Mutual recognition of an authorisation of a BP family	For one biocidal product type and one user type : 30 000€ For each additional user type : + 6000€ For each additional product type : + 6000€ For each biocidal product in the family : + 800€/product
Union authorisation of a BP	For one biocidal product type and one user type : 40 000€ For each additional user type : + 8000€ For each additional product type : + 8000€ When the product contain at least one AS candidate to substitution : + 10 000 €
Union authorisation of a BP family	For one biocidal product type and one user type: 80 000€ For each additional user type: + 16 000€ For each additional product type: + 16 000€ When the product contain at least one AS candidate to substitution: + 20 000 € For each biocidal product in the family: + 800€/product
Annual fee	None
Other (please specify)	Transformation of a provisional MA into a MA for a BP : 5000€ Transformation of a provisional MA into a MA for a BP family : 10 000€ For each biocidal product in the family : +800€/product Provisional authorisation for a BP which is the reference product for a new AS - if FR is not EMR : 40 000€; if FR is EMR : none Provisional authorisation for a BP family which is the reference product for a new AS - if FR is not EMR : 80 000€; if FR is EMR : none For each biocidal product in the family : +800€/product

2.3. Measures in favour of small and medium enterprises (SMEs)

Do you have specific national measures or legislation favouring SMEs?								
	Do١	vou have	specific nation	nal measures	or le	nislation	favouring	SMFs?

O Yes

No

2.4. Non-compliance and penalties

Do you have specific national measures or legislation concerning non-compliance and penalties applicable for infringements on the implementation of the BPR?

Yes

O No

Please specify below the national regulation(s) and refer to the corresponding link to the relevant website with the requested information. If available in English please include the link to the English version

The are two types of measures, all described in the Environmental Code : - administrative measures - penal measures. Administrative measures : - Article L521-17 : possibility for the national enforcement authority to start a formal notice procedure, with a specific perio d, and require the compliance to the BPR. https://www.legifrance.gouv.fr/affichCodeArticle.do?idArticle=LEGIARTI000022964116&cidTexte=LEGITEXT000006074220&dateTexte=201010 23 - Article L521-18 (point 1) : In case of non-compliance after the formal notice procedure, the NEA has the possibility to order a fine payment (up to 15000 euros) and a daily penalty payment as long as the infringement persists (up to 1500 euros per day). https://www.legifrance.gouv.fr/affichCodeArticle.do;jsessionid=904F85F50756DF99C64D484AE63EA5B9.tplgfr38s 2?idArticle=LEGIARTI000 - Article L521-18 (point 2 and 3) : Possibility to order a ban on the placing on the market of substances, biocidal products and treated articles, possibility to order the importer to return the substance, product or treated article outside the EU. https://www.legifrance.gouv.fr/affichCodeArticle.do;jsessionid=904F85F50756DF99C64D484AE63EA5B9.tplgfr38s 2?idArticle=LEGIARTI000 - Article L522-15 : In case of non-compliance after a formal notice procedure, the NEA has the possibilty to take measures in ord er to stop the use of a non-compliant substance, product or treated article. The responsible of the placing on the market could b e asked to ensure their recovery and their disposal. https://www.legifrance.gouv.fr/affichCodeArticle.do?cidTexte=LEGITEXT000006674220&idArticle=LEGIARTI000006834394&dateTexte=&categ orieLien=cid Penal measures : Several penal measures are provided by law in the Environmental Code, depending on the nature of the non-compliance. - Article L.522-16 (point I) : the placing on the market of a non-compliant active substance, biocidal product or treated article is punishable by imprisonment of two years and a fine of 75000 euros https://www.legifrance.gouv.fr/affichCodeArticle.do?idArticle=LEGIARTI000031559456&cidTexte=LEGITEXT000006074220&dateTexte=201512 - Article L.522-16 (point II) : the use of a biocidal product without complying with its authorisation is punishable by imprisonm ent of 6 months and a fine of 7500 euros https://www.legifrance.gouv.fr/affichCodeArticle.do?idArticle=LEGIARTI000031559456&cidTexte=LEGITEXT000006074220&dateTexte=201512

2.5. Imported treated articles

Do you have specific national measures or legislation that regulates whether imported treated articles contain only approved active substances?

incomplete declarations, non-compliant labelling for a biocidal product or a treated article...

- Yes
- O No

Please specify below the national regulation(s) and refer to the corresponding link of the relevant website with the requested information. If available in English please include the link to the English version

- Article R522-25 : various penalties are specified, in case of absence of declaration of the product on the national databases,

Environmental Code - article L.522-16
https://www.legifrance.gouv.fr/affichCodeArticle.do?idArticle=LEGIARTI000031559456&cidTexte=LEGITEXT000006074220&dateTexte=201512
04

3. Placing and making available on the market of biocidal products

3.1. Authorisation procedures

3.1.1. Authorisations

Please indicate in the tables below the figures related to the various procedures since the entry into application of the BPR (1st September 2013)

3.1.1.a. National authorisations

	2013	2014	2015	2016	2017	2018	2019	Total number
Authorisations granted on the basis of Article 19(5)	105	138	37	42	9	8	17	356
Provisional authorisations granted for products containing new active substances (Article 55(2))	1	0	0	0	0	0	0	1

3.1.1.b. Mutual recognitions - concerned Member State

	2013	2014	2015	2016	2017	2018	2019	Total number
Derogations (Article 37)	0	1	8	2	5	9	4	29

3.1.1.c. Authorisations of products containing active substances meeting exclusion criteria (Article 5.2)

	2013	2014	2015	2016	2017	2018	2019	Total number
Total number of applications assessed	100	171	37	42	30	9	17	391
Number of products authorised (conditions met for all or some of the uses)	99	137	36	42	27	9	17	352
Number of products not authorised (conditions not met for any of the uses)	1	34	1	0	3	0	0	9

3.1.1.d. Comparative assessments (Article 23)

	2013	2014	2015	2016	2017	2018	2019	Total number
Total number of applications evaluated	60	115	31	52	66	15	25	364
Number of applications evaluated resulting in a granted authorisation without restrictions	59	80	30	51	61	15	25	321
Number of applications evaluated resulting in restrictions (Article 23(3))	0	0	0	0	0	0	0	0
Number of applications evaluated resulting in restrictions (Article 23(3))	1	35	1	1	5	0	0	43

3.1.2. Renewal of authorisations

Please indicate in the tables below the figures related to the renewals of authorisations for making available on the market of biocidal products

3.1.2.a. National authorisations

	2013	2014	2015	2016	2017	2018	2019	Total number
Authorisations granted on the basis of Article 19(5)	0	0	0	0	7	94	22	123
Provisional authorisations granted for products containing new active substances (Article 55(2))	0	0	0	0	0	0	0	0

3.1.2.b. Mutual recognitions - concerned MS

	2013	2014	2015	2016	2017	2018	2019	Total number
Derogations (Article 37)	0	0	0	0	0	0	0	0

3.1.2.c. Authorisations of products containing active substances meeting exclusion criteria (Article 5.2)

	2013	2014	2015	2016	2017	2018	2019	Total number
Total number of applications assessed	0	0	0	0	7	94	22	123
Number of products authorised (conditions met for all or some of the uses)	0	0	0	0	7	93	22	122
Number of products not authorised (conditions not met for any of the uses)	0	0	0	0	0	1	0	1

3.1.2.d. Comparative assessments (Article 23)

	2013	2014	2015	2016	2017	2018	2019	Total number
Total number of applications evaluated	0	0	0	0	4	70	16	90
Number of applications evaluated resulting in a granted authorisation without restrictions	0	0	0	0	0	19	16	35
Number of applications evaluated resulting in restrictions (Article 23(3))	0	0	0	0	4	51	0	55
Number of applications evaluated resulting in restrictions (Article 23(3))	0	0	0	0	0	0	0	0

3.2. Other BPR procedures for biocidal products

The BPR contains specific procedures that allow the making available of the market of products without an authorisation. Please indicate the related information in the tables below.

3.2.a. Derogations pursuant to Article 55(1)

	2013	2014	2015	2016	2017	2018	2019	Total number
Number of requests received	0	1	0	1	2	2	3	9
Permits granted	0	1	0	1	2	1	1	6
Permits not granted	0	0	0	0	0	1	2	3

3.2.b. Research and development (Article 56)

	2013	2014	2015	2016	2017	2018	2019	Total number
Number of notifications received	0	1	0	4	6	4	5	20
Number of prohibitions	0	0	0	0	0	0	0	0

3.3. Number of biocidal products made available on the Member State market authorised under the transitional measures (Article 89)

Please provide the information available, per main group of product-types, on biocidal products made available on the market authorised under transitional measures

	2013	2014	2015	2016	2017	2018	2018	Total number
Main group 1 Disinfectants (PT1 - PT5)	25	35	33	0	0	0	0	93
Main group 2 Preservatives (PT6 - PT13)	0	0	0	0	0	0	0	0
Main group 3 Pest control (PT14 - PT20)	9	7	9	0	0	0	0	25
Main group 4 Other biocidal products (PT21 - PT22)	0	0	0	0	0	0	0	0

4. Information on enforcement activities

4.1. BPR enforcement strategy

Has an overall strategy been implemented in the Member State for the enforcement of the BPR?

Yes

O No

Please describe it and, if it is publicly available, provide the corresponding link. If available in English please include the link to the English version

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An annual DGCCRF biocide control plan has been in place since 2006.
This annual plan includes the control of detergents since 2017.
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4.2. Control system in the Member States and results of official controls

Please give a brief overview of the way official controls are carried out in your Member State, with special emphasis on the following processes:

- making available on the market of biocidal products;
- use of biocidal products;
- placing on the market of treated articles

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DGGCRF is responsible for the control of biocidal products for consumers.

Treated articles were subject to a specific control plan in 2019 under the BEF1.
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4.3. Enforcement authorities involved in official controls

Please provide the denomination of the enforcement authorities involved in official controls. If applicable, please also provide the links to the relevant websites

	BPR enforcement authority(ies) involved	Website
Controls on placing and making biocidal products available on the market	DGCCRF for consumer products	https://www.economie.gouv.fr/dgccrf
Controls on placing on the market of treated articles	DGCCRF for consumer products	https://www.economie.gouv.fr/dgccrf

4.3.1. Complaints

Have there been complaints (information about suspected infringements of the BPR rules) received by enforcement authorities in relation to the implementation of the BPR?

	2013	2014	2015	2016	2017	2018	2019	Total number
Number of complaints								

4.4. Controls addressing different parts of the supply chain

The reporting Member State is requested to provide information on the official controls on the implementation of the BPR that have been performed since the entry into force of the Regulation.

Please provide below the information available on the official controls performed with a focus on the following information, where available: number of controls performed, resources used, outcome of controls (in terms of number and type of non-compliances)

4.4.1. Official controls on compliance with BPR rules for making available on the market of biocidal products

• (Total number of) controls on biocidal products made available on the market / (Number of) illegal products made available and points of non-compliance

If detailed figures are available please provide them in the table below

Controls on biocidal products - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants	699	196	855	240	773	209	956	315	894	419	1162	476	826	226
MG 2 Preservatives	148	41	155	43	201	54	175	58	129	61	99	40	200	55
MG 3 Pest control	420	118	1004	281	1272	344	1137	375	793	372	1160	477	1598	438
MG 4 Other biocidal	32	_	0	_	_	4	20	40	00	40	40	20	00	20
products	32	9	9	2	5	1	29	10	28	13	49	20	80	22

· Controls on records kept by authorisation holders, in accordance with article 68 of the BPR

If detailed figures are available please provide them in the table below

Controls on records kept by authorisation holders - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants														
MG 2 Preservatives														
MG 3 Pest control														
MG 4 Other biocidal														
products														

• Controls on the classification, packaging and labelling of biocidal products (article 69 of the BPR, and Regulation 1272/2008 on the classification, labelling and packaging of substances and mixtures (CLP))

If detailed figures are available please provide them in the table below

Controls on classification, packaging and labelling of biocidal products - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants	699	126	855	162	773	186	956	172	894	185	1162	279	826	120
MG 2 Preservatives	148	27	155	29	201	48	175	31	129	27	99	24	200	29
MG 3 Pest control	420	75	1004	191	1272	305	1137	205	793	164	1160	278	1598	232
MG 4 Other biocidal products	32	6	9	2	5	1	29	5	28	6	49	11	80	12

• Controls on safety data sheets (article 70 of the BPR, and article 31 of Regulation 1907/2006 on the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH))

If detailed figures are available please provide them in the table below

Controls on safety data sheets - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants	699	??	855	9	773	23	956	??	894	49	1162	29	826	31
MG 2 Preservatives	148	??	155	2	201	6	175	??	129	7	99	2	200	8
MG 3 Pest control	420	??	1004	10	1272	38	1137	??	793	44	1160	30	1598	61
MG 4 Other biocidal products	32	??	9	1	5	0	29	??	28	1	49	1	80	3

• Controls on advertisement of biocidal products (article 72 of the BPR and CLP)

If detailed figures are available please provide them in the table below

Controls on advertisment of biocidal products - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants	699	126	855	231	773	116	956	220	894	97	1162	186	826	70
MG 2 Preservatives	148	27	155	42	201	30	175	40	129	14	99	15	200	17
MG 3 Pest control	420	75	1004	271	1272	191	1137	262	793	85	1160	186	1598	136
MG 4 Other biocidal	32	6	9	2	5	1	29	7	28	3	49	8	80	7
products				_		ļ ·								

• Controls on the inclusion of active substance suppliers in the official list (article 95(2) of the BPR)

If detailed figures are available please provide them in the table below

Controls on the inclusion of active substance suppliers in the Article 95 list - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants											3	0	5	0
MG 2 Preservatives											2	0	0	0
MG 3 Pest control											2	0	5	0
MG 4 Other biocidal											0	_	0	0
products											0	0	0	U

4.4.2. Official controls on biocidal products made available on the market during the transitional period

• Controls to ensure that the biocidal products on the market contain active substances included in the review programme (Article 89(2) of the BPR)

If detailed figures are available please provide them in the table below

Controls to ensure that the biocidal products on the market contain active substances included in the review programme - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants	699	56	855	51	773	31	956	16	894	25	1162	14	826	7
MG 2 Preservatives	148	12	155	9	201	8	175	4	129	4	99	1	200	2
MG 3 Pest control	420	33	1004	60	1272	51	1137	23	793	23	1160	14	1598	16
MG 4 Other biocidal products	32	2	9	0	5	0	29	0	28	1	49	0	80	1

• Controls on the inclusion of active substances suppliers in the official list (article 95(2) of the BPR)

If detailed figures are available please provide them in the table below

Controls on the inclusion of active substance suppliers in the Article 95 list - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants											3	0	5	0
MG 2 Preservatives											2	0	0	0
MG 3 Pest control											2	0	5	0
MG 4 Other biocidal											0	0	0	0
products											0	U	0	U

• Controls on compliance of the biocidal products made available on the market with national legislation (where relevant)

 ${\tt SIMMBAD} \ \ {\tt declaration} \ \ ({\tt inventory} \ \ {\tt and} \ \ {\tt annual} \ \ {\tt quantities} \ \ {\tt sold})$

If detailed figures are available please provide them in the table below

Controls on compliance of the biocidal products made available with national legislation - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants	699	98	855	111	773	116	956	134	894	133	1162	139	826	47
MG 2 Preservatives	148	21	155	20	201	30	175	24	129	19	99	12	200	11
MG 3 Pest control	420	59	1004	131	1272	191	1137	159	793	118	1160	139	1598	91
MG 4 Other biocidal products	32	4	9	2	5	1	29	4	28	4	49	6	80	4

4.4.3. Official controls on manufacturers

• Controls regarding the availability of the appropriate documentation in relation to the manufacturing process, as indicated in article 65 (2) of the BPR

If detailed figures are available please provide them in the table below

Controls regarding the availability of the appropriate documentation related to the manufacturing process - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants														
MG 2 Preservatives														
MG 3 Pest control														
MG 4 Other biocidal														
products														

4.4.4. Official controls on end-users and residues

• Controls regarding the use of the biocidal products according to the terms and conditions of the authorisation, as stipulated in article 17(5) of the BPR

If detailed figures are available please provide them in the table below

Controls regarding the use of the biocidal products according to the terms and conditions of the authorisation - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants														
MG 2 Preservatives														
MG 3 Pest control														
MG 4 Other biocidal														
products														

• Controls on residue levels of active substances in food and feed (PT3, 4, 5, 18, 19 and 21)

If detailed figures are available please provide them in the table below

Controls on residue levels of active substances in food and feed - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
PT 3														
PT 4														
PT														
18														
PT														
19														
PT														
21														

4.4.5. Official controls on treated articles

• Controls concerning the active substance(s) present in the treated articles (articles 58(2) and 94 of the BPR)

A specific control plan DGCCRF to treated articles in 2019 under the ${\tt BEF1}$

If detailed figures are available please provide them in the table below

Controls concerning the active substance(s) present in the treated articles - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants	0	0	0	0	15	??	32	2	25	2	20	1	22	7
MG 2 Preservatives	0	0	0	0	0	0	0	0	6	??	10	4	59	2
MG 3 Pest control	0	0	0	0	0	0	8	0	7	??	15	2	87	4
MG 4 Other biocidal products	0	0	0	0	0	0	0	0	0	0	0	0	0	0

• Controls on the correct labelling of the treated articles (article 58 of the BPR)

A specific control plan DGCCRF to treated articles in 2019 under the ${\tt BEF1}$

If detailed figures are available please provide them in the table below

Controls on the correct labelling of the treated articles - Total number of controls per year and non-compliances (NC) identified

	Total	NC												
	2013	2013	2014	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
MG 1 Disinfectants	0	0	0	0	15	7	32	15	25	5	20	1	12	6
MG 2 Preservatives	0	0	0	0	0	0	0	0	6	2	10	6	44	19
MG 3 Pest control	0	0	0	0	0	0	8	8	7	2	15	4	63	25
MG 4 Other biocidal	0	0	0	0	0	0	0	0	0	0	0	0	0	0
products	0	0	0	0	U	U	0	0	0	0	0	0	0	0

5. Poisoning incidents

5.1. Poisonings involving biocidal products, severity of the impact

Please provide below an overview of the information reported since the entry into operation of BPR on poisoning incidents involving biocidal products, indicating the active substances and product-types most frequently involved and those involved in incidents resulting in severe health impairments or death (fatal or near fatal incidents)

No overall overview is available.

Poison centres are currently updating their formulation database so that biocidal products can be automatically identified among all chemicals. This update will allow a systematic identification of incidents in which biocidal products are involved, by PTs. This work is still on-going. It has been delayed due to coronavirus crisis.

In the past 7 years several report dedicated to specific topics have been made. Studies in which poisoning incidents involving biocidal products between 2013 and 2019 were reported are listed elow:

Adverse events collected during vector control campaigns using deltamethrin or Bti (Bacillus thuringiensis israelensis), France, 2 006-2013. December 2013

- http://www.centres-antipoison.net/cctv/Rapport CCTV Rapport Lutte antivectorielle vf.pdf

Exposure to products containing phosphides - Retrospective study of incidents recorded by the French poison and toxicovigilance c entres (1999-2013). September 2015

http://www.centres-antipoison.net/cctv/CCTV Rapport phosphures D111.pdf

Exposure to products containing TP8 biocides: Cases registered by French poison control centers between 2012 and 2014. September 2016

No public version available on a website

Exposure to products containing phosphides as part of a harbour activity or when opening a container - Retrospective study of inc idents recorded by the French poison and toxicovigilance centres (1999-2017). September 2018

- https://www.anses.fr/fr/system/files/Toxicovigilance2018SA0290Ra.pdf

Accidental exposure to chlorinated cleaning products for swimming pools and spas - Retrospective study of incidents recorded by the French poison and toxicovigilance centres (2010-2019). June 2019

https://www.anses.fr/fr/system/files/Toxicovigilance2019SA0192Ra.pdf

Skin and eye reactions to insect repellent bracelets- Retrospective study of the incidents recorded by the French poison and toxi covigilance centres (2012-2019). November 2019

https://www.anses.fr/fr/system/files/Toxicovigilance2012SA0277Ra.pdf
 No overall overview is available.

Poison centres are currently updating their formulation database so that biocidal products can be automatically identified among all chemicals. This update will allow a systematic identification of incidents in which biocidal products are involved, by PTs. This work is still on-going. It has been delayed due to coronavirus crisis.

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Exposure to products containing phosphides - Retrospective study of incidents recorded by the French poison and toxicovigilance c entres (1999-2013). September 2015

http://www.centres-antipoison.net/cctv/CCTV_Rapport_phosphures_D111.pdf

Exposure to products containing TP8 biocides: Cases registered by French poison control centers between 2012 and 2014. September 2016

No public version available on a website

Exposure to products containing phosphides as part of a harbour activity or when opening a container - Retrospective study of inc idents recorded by the French poison and toxicovigilance centres (1999-2017). September 2018

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https://www.anses.fr/fr/system/files/Toxicovigilance2012SA0277Ra.pdf

If detailed figures are available please provide them in the table below

Number of poisoning incidents related to biocidal products per year by poisoning severity

	2013 Fatal/near fatal	2013 Other	2014 Fatal/near fatal	2014 Other	2015 Fatal/near fatal	2015 Other	2016 Fatal/near fatal	2016 Other	2017 Fatal/near fatal	2017 Other	2018 Fatal/near fatal	2018 Other	2019 Fatal/near fatal	2019 Other
MG 1														
Disinfectants														
MG 2														
Preservatives														
MG 3 Pest														
control														
MG 4 Other														
biocidal products														

6. Helpdesk functioning

Please fill in in the following table the information regarding the number of enquiries that Helpdesks receive per year.

Note: if your system does not differentiate the queries according to their topic (active substances, biocidal products, treated articles) please indicate the total number of queries per year in the last row.

	2013	2014	2015	2016	2017	2018	2019
Number of enquiries on active substances	No data available before 2018						
Number of enquiries on biocidal products							
Number of enquiries on treated articles							
Total number of enquiries per year						870	800

6.1. Advice to small and medium-sized enterprises (SMEs)

Following Article 81(2) of the BPR, competent authorities have to provide advice to the applicants and in particular to SMEs.

Does your Member State provide specific adv	ce to SMEs?	
○ Yes		
No		

7. Sustainable use measures

In accordance with Article 18 of the BPR, a Commission Report on the sustainable use of biocidal products was submitted to the European Parliament and the Council in 2016, compiling the information provided by Member States. Please find below some questions that are requested in order to follow-up on this

7.1. Availability of Best Practices Documents in the Member States

Are Best Practices Documents used or developed for reducing the use of biocidal products to a minimum or for using biocides with less impact on human health and the
environment?
○ Yes
○ No

7.2. Availability of certifications or training schemes for professional users

Ar ?

re certification procedures or training schemes in place (organised by e.g. eCAs, public authorities, sector organisations) for professional us	ers of biocidal products
Yes	
O No	
O Not anymore	

Please specify which kind of biocidal products or applications are covered by those schemes and include the corresponding links of the relevant websites with information

	Biocidal products or applications covered	Name of the certification or training scheme	Year
1	The products concerned by this certification are—some TP2,3 and 4 and all TP8, 14 15, 18 and 20 (only professional products). All professional who buy, sell or use these products need a certification except if the product is used in a transformation / production cycle. Firefighters do not need "certibiocide" as soon as they intervene in emergency situations. The link to the national decree https://www.legifrance.gouv.fr/affichTexte.do?cidTexte=JORFTEXT000028214219&dateTexte=20200617	certibiocide	this certification was created in 2015
2			
3			

7.3. Information to the public

Have measures been taken to provide the public with appropriate information about benefits and risks associated with biocidal products and ways of minimisi	ing their
use? (Article 17(5) of the BPR)	

YesNo

Please specify which kind of information is available in your Member State (e.g. information campaigns, regulatory measures) and include the corresponding links of the relevant websites with information.

	Type of measure	Year	Details
1	Egalim decree regulatory measures concerning prohibition of self service sale to non professional users of certain categories of biocidal products	2019	https://www.legifrance.gouv.fr/affichTexte.do? cidTexte=JORFTEXT000039223395&categorieLien=id
2	Egalim decree regulatory measures concerning prohibited commercial practices for certain categories of biocidal products	2019	https://www.legifrance.gouv.fr/affichTexte.do? cidTexte=JORFTEXT000038689025&categorieLien=id
3	Egalim decree regulatory measures concerning commercial advertising for certain categories of biocidal products	2019	https://www.legifrance.gouv.fr/affichTexte.do? cidTexte=JORFTEXT000038689038&categorieLien=id
4			

7.4. Measures to address the risk related to the use of biocidal products

Have measures been taken to address the risks related to the use of biocidal products in specific areas such as schools, workplaces, kindergartens or public spaces?

Yes

O No

Please specify which kind of information is available in your Member State (e.g. information campaigns, regulatory measures) and refer to the corresponding links of the relevant websites with information.

	Type of measure	Year	Area covered	Details
	Egalim decree regulatory measures concerning prohibition of self service sale to non professional users of certain categories of biocidal products	2019		
	2 Egalim decree regulatory measures concerning prohibited commercial practices for certain categories of bioci products	dal 2019		
	3 Egalim decree regulatory measures concerning commercial advertising for certain categories of biocidal products	2019		
-	4			

8. Nanomaterials

Please provide information regarding the use of nanomaterials in biocidal products (Articles 19 and 69 of the BPR), per main group of product-types

MG 1 Disinfectants

	Product name	Nanomaterial	Brief explanations	Safety measures (Yes/No)	Year
1					
2					
3					
4					

MG 2 Preservatives

	Product name	Nanomaterial	Brief explanations	Safety measures (Yes/No)	Year
1					
2					
3					
4					

MG 3 Pest control

	Product name	Nanomaterial	Brief explanations	Safety measures (Yes/No)	Year
1	MITE KILLER	Silicium dioxide	Biocidal active substance	NO	2019
2					
3					
4					

MG 4 Other biocidal products

	Product name	Nanomaterial	Brief explanations	Safety measures (Yes/No)	Year
1					
2					
3					
4					

9. Any other comment

ANSES : Seuls les produits contenant des substances actives qui sont des nanomateriaux ont été indiqués. Nous n'avons pas l'information pour les autres substances entrant dans la composition des produits.

Contact

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