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BPR Article 65(3) reporting

1. General information

1.1. Member State

For which Memeber State* are you reporting?

(*In this survey "Member State" refers to EU Member States, Iceland, Liechtenstein, Norway and Switzerland)

- ○ Austria
- ○ Belgium
- ○ Bulgaria
- ○ Croatia
- ○ Cyprus
- ○ Czech Republic
- ○ Denmark
- ○ Estonia
- ○ Finland
- ○ France
- ○ Germany
- ○ Greece
- ○ Hungary
- ○ Iceland
- ○ Ireland
- ○ Italy
- ○ Latvia
- ○ Liechtenstein
- ○ Lithuania
- ○ Luxembourg
- ○ Malta
- ○ Netherlands
- ○ Norway
- ○ 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)
	[redacted]	Chemicals Office of the Republic of Slovenia (CORS), Ministry of Health	[redacted]	

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.

one

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	Chemicals Office of the Republic of Slovenia	www.uk.gov.si
Advice to applicants/helpdesks on biocidal products	Chemicals Office of the Republic of Slovenia	www.uk.gov.si
Advice to applicants/helpdesks on treated articles	Chemicals Office of the Republic of Slovenia	www.uk.gov.si

Assessment of active substances	Chemicals Office of the Republic of Slovenia	www uk.gov si
Assessment and authorisation of biocidal products	Chemicals Office of the Republic of Slovenia	www uk.gov si
Other (e.g. authority in charge of setting up the whole organisational framework for the BPR implementation, of adopting national legislation)	Chemicals Office of the Republic of Slovenia	www uk.gov si

1.3.3. Other bodies involved in the implementation of the BPR

	Authority/organisation involved	Website
Poison centre	Centre for Clinical Toxicology and Pharmacology	https://www kclj si/index.php?dir=/pacienti_in_obiskovalci/klinike_in_oddelki/interna_klinika/center_za_klinicno_toksikologijo_in_farmakologijo
Animal poison centre	/	
Other	National Institute of Public Health, Agricultural Institute of Slovenia	https://www nijz.si/en, https //www kis.si/en/

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
- ( ) 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.

Decree implementing Regulations (EU) concerning making available on the market and use of biocidal products, OJ RS No 81/2018 regulates conditions for the making available of biocidal products in the Republic of Slovenia during transitional period (Article 5 - notification approval procedure), competent authority, supervision, offences and sanctions for the purpose of implementing Regulation (EU) No 528/2012. Annex to this Decree regulates fees for services provided by the competent authority in connection with the procedures under Regulation (EU) No 528/2012 and Decree RS No 81/18.

National transitional measures for making biocidal products available (notification) as regulated in decree:

"Article 5
(1) An application for notification of a biocidal product that contains an active substance referred to in the first subparagraph of Article 89(2) of Regulation (EU) No 528/2012 shall be submitted no later than the date of inclusion of the active substance in the Union list of approved active substances.
(2) Biocidal products shall be available in the territory of the Republic of Slovenia after they have been entered into the national register by the Office.
(3) Where a biocidal product referred to in paragraph one of this Article is used in whole for the manufacture of a new biocidal product, only the final product shall be notified.
(4) The application form for notification, change in notification or completion of notification of a biocidal product shall be published on the website of the Office.
(5) In the notification procedure, the Office shall verify compliance of a biocidal product based on the following:
- fulfilment of conditions for the active substance referred to in paragraph one of this Article;
- combination of the active substance and product type; and
- origin and supply of active substances under Article 95 of Regulation (EU) No 528/2012.
(6) In the notification procedure, or in respect of a biocidal product that has already been notified, the Office may verify the following:
- the chemical, physical and technical properties of the product;
- information relating to the ecotoxicity and toxicity of the biocidal product and its efficacy;
- the product's presentation and appearance;
- the method of allowing the product to be made available; and
- labelling and instructions for use.
(7) If, on the basis of the information submitted, the Office considers that the intended use of the biocidal product would constitute an unacceptable risk to humans, animals or the environment, it may require the applicant to provide additional explanatory information and propose additional measures for the safe use of the biocidal product.
(8) When it is not possible to ensure safe use of the biocidal product, the Office shall refuse the notification, change the entry or remove the notified biocidal product from the national register.
(9) If, following the risk assessment, the active substance in the notified biocidal product has not been included in the Union list of approved active substances, or if an application for the issue of a national authorisation or Union authorisation for the biocidal product has not been submitted by the date of inclusion of the active substance in the Union list of approved active substances, in the national register, the Office shall specify the date until which the notified biocidal product may be available on the market, in accordance with Regulation (EU) No 528/2012, and shall remove the biocidal product from the national register after the expiry of that time limit."

Regulation is available in Slovene language only on the following website: http://www.pisrs.si/Pis.web/pregledPredpisa?id=URED7799# Unofficial translation is available by the CA.

General information on biocides is available on the website: https://www.gov.si/teme/biocidni-proizvodi/

Information on transitional notification procedure for companies and general public is available on: http://evem.gov.si/info/dovoljenja/dejavnost/13738/dovoljenje/12493/prikaziDovoljenje/

During the reporting period, legislation on biocidal products changed twice. Act Repealing the Biocidal Products Act, OJ RS, No 25/14 of 2.4.2014 with secondary legislation and Decree implementing Regulations (EU) concerning the making available on the market and use of biocidal products, OJ RS No 20/14 have been replaced by above mentioned Decree of RS No 81/2018.
National transitional approval procedure was in place since 2000.

2.2. Applicable fees

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

- (●) Yes
- ( ) 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

Anex of the Decree implementing Regulations (EU) concerning making available on the market and use of biocidal products, OJ RS No. 81/2018 regulates amounts of fees. National measures are regulated by Article 6:

"(1) For the costs relating to procedures for approving active substances and granting authorisations for biocidal products referred to in Regulation (EU) No 528/2012, for active substances and biocidal products for which the Office is responsible, the applicant shall pay the fee set out in Annex, which is an integral part of this Decree.

(2) After receipt of an application referred to in paragraph one of Article 3 of this Decree, the Office shall set the initial fee amount depending on the estimated extent and complexity of the procedure and shall inform the applicant of the amount and method of payment of the fee and the deadline for same.

(3) Before the completion of the procedure, the Office shall evaluate the actual work performed and shall invite the applicant to pay an additional amount of fee or shall refund them the overpaid amount of fee referred to in the preceding paragraph.

(4) Fees shall be paid into the general government revenue sub-account, in accordance with the regulation governing sub-accounts and the manner of payment of compulsory charges and other general government revenue.

(5) In accordance with Article 80 of Regulation (EU) No 528/2012, from the fees paid under points 1 and 2 of Annex to this Decree, appropriate funds shall, as a priority, be earmarked in the financial plan of the Office to cover the entire costs of the phases of the procedure referred to in points 1.B. in 2.1.B. of Annex to this Decree and the related material and technical conditions and administrative and technical support.

(6) If the applicant withdraws their application at any stage of the procedure, the costs of the procedure shall be charged in proportion to the work performed by the Office. Costs shall also be charged in the same manner in cases where the Office accepts to undertake procedures that have been only partially completed by other competent authorities."

Regulation is available in Slovene language only on the following link: http://www.pisrs.si/Pis.web/pregledPredpisa?id=URED7799#

Annex includes fees for the approval of active substances with description of the procedure and specification of costs referred according to different phases of the evaluation procedure and fees for granting authorisations for biocidal products with description of the procedure and specification of costs according to different phase of the evaluation procedure. Fees are established for a single biocidal product and a biocidal product family for procedures in which CORS acts as a reference member state/evaluating competent authority or as concerned member state/competent authority. Fee for notification of biocidal products according transitional procedure is also regulated (210 EUR).
All fees are in euros.

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	chemical: 28,500 - 117,000; microorganism: 85,600
Evaluation of an active substance for Annex I inclusion	58,500
Authorisation of a biocidal product (BP)	23,400
Authorisation of a BP family	46,800
Mutual recognition of an authorisation of a BP	2,600 - 3,000
Mutual recognition of an authorisation of a BP family	5,200 - 6,000
Union authorisation of a BP	32,760
Union authorisation of a BP family	65,520
Annual fee	0
Other (please specify)	see Annex to the Decree on implementation of BPR, RS No 81/18

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

Do you have specific national measures or legislation favouring SMEs?

- ○ 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
- ○ 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

Decree implementing Regulations (EU) concerning making available on the market and use of biocidal products, OJ RS No. 81/2018 regulates specific national measures concerning non-compliance and penalties for infringements on the implementation of the BPR. National measures are regulated by Articles 8 and 9:

"Article 8

(1) A legal person shall be fined from EUR 5,000 to EUR 30,000 for the following offences:
1. the making available on the market and for use of a biocidal product in contravention of Article 5 of this Decree;
2. the making available on the market and for use of a biocidal product that is not authorised in accordance with Article 17 of Regulation (EU) No 528/2012;
3. failure to notify the Office of any unexpected or adverse effects in accordance with Article 47(1) of Regulation (EU) No 528/2012;
4. the making available of a biocidal product for use by users for whom the making available and use of the biocidal product have not been authorised;
5. failure to present, as the supplier of an active substance or manufacturer of a biocidal product, evidence of compliance of active substances with Article 95 of Regulation (EU) No 528/2012 or failure to present such evidence within the time limit specified by a chemicals inspector;
6. failure to meet the requirements and conditions for use by professional users or trained professional users in accordance with paragraph two or three of Article 4 of this Decree.

(2) A sole trader or a self-employed person shall be fined from EUR 5,000 to EUR 15,000 for an offence referred to in the preceding paragraph.

(3) The responsible person of a legal person or the responsible person of a self-employed person shall be fined from EUR 2,000 to EUR 5,000 for an offence referred to in paragraph one of this Article.

Article 9

(1) A legal person shall be fined from EUR 2,500 to EUR 15,000 for the following offences:
1. the use of a biocidal product in contravention of Article 17(1) and (5) of Regulation (EU) No 528/2012;
2. the making available on the market and for use of a biocidal product contrary to the parallel trade permit, as laid down in Article 53 of Regulation (EU) No 528/2012;
3. the making available on the market and for use of a biocidal product contrary to the emergency authorisation, provisional authorisation or exceptional authorisation granted in accordance with Article 55 of Regulation (EU) No 528/2012;
4. the making available on the market and for use of a biocidal product contrary to the authorisation for the purposes of research or development granted in accordance with Article 56 of Regulation (EU) No 528/2012;
5. the making available on the market and for use of a treated article in contravention of Article 58(2) of Regulation (EU) No 528/2012;
6. the making available on the market and for use of a treated article that is not labelled in accordance with Article 58(3) of Regulation (EU) No 528/2012;
7. failure to keep records of biocidal products in accordance with Article 68(1) of Regulation 528/2012/EU;
8. the making available on the market and for use of a biocidal product which is not classified, packaged or labelled in accordance with Article 69(1) of Regulation (EU) No 528/2012 or for which no safety data sheet has been prepared in accordance with Article 70 of Regulation (EU) No 528/2012;
9. the advertising of a biocidal product in contravention of Article 72 of Regulation (EU) No 528/2012.

(2) A sole trader or a self-employed person shall be fined from EUR 2,500 to EUR 10,000 for an offence referred to in the preceding paragraph.

(3) The responsible person of a legal person, the responsible person of a sole trader or the responsible person of a self-employed person shall be fined from EUR 1,500 to EUR 3,000 for an offence referred to in paragraph one of this Article."

Regulation is available in Slovene language only on the following link: http://www.pisrs.si/Pis.web/pregledPredpisa?id=URED7799#

2.5. Imported treated articles

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

- ○ Yes
- ◉ No

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)	0	0	0	0	0	0	0	0
Provisional authorisations granted for products containing new active substances (Article 55(2))	0	0	0	0	0	0	0	0

3.1.1.b. Mutual recognitions - concerned Member State

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

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	0	0	5	18	11	2	2	38
Number of products authorised (conditions met for all or some of the uses)	0	0	5	17	11	2	2	37
Number of products not authorised (conditions not met for any of the uses)	0	0	0	1	0	0	0	1

3.1.1.d. Comparative assessments (Article 23)

	2013	2014	2015	2016	2017	2018	2019	Total number
Total number of applications evaluated	0	0	0	0	3	2	0	5
Number of applications evaluated resulting in a granted authorisation without restrictions	0	0	0	0	0	2	0	2
Number of applications evaluated resulting in restrictions (Article 23(3))	0	0	0	0	3	0	0	3
Number of applications evaluated resulting in restrictions (Article 23(3))								0 prohibitions

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	0	0	0	0
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	1	0	0	21	7	29
Number of products authorised (conditions met for all or some of the uses)	0	0	1	0	0	21	7	29
Number of products not authorised (conditions not met for any of the uses)	0	0	0	0	0	0	0	0

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	0	0	0	0
Number of applications evaluated resulting in a granted authorisation without restrictions	0	0	0	0	0	0	0	0
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))	0	0	0	0	0	0	0	0 prohibitions

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	0	0	0	0	0	0	0
Permits granted	0	0	0	0	0	0	0	0
Permits not granted	0	0	0	0	0	0	0	0

3.2.b. Research and development (Article 56)

	2013	2014	2015	2016	2017	2018	2019	Total number
Number of notifications received	0	0	0	0	0	0	0	0
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)	30	54	123	153	130	107	137	734
Main group 2 Preservatives (PT6 - PT13)	7	31	31	55	55	38	45	262
Main group 3 Pest control (PT14 - PT20)	7	38	38	54	54	35	50	276
Main group 4 Other biocidal products (PT21 - PT22)	0	4	1	36	36	3	2	82

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
- ( ) 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

No

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

all mentioned issues are covered

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	Chemicals Office of the Republic of Slovenia	https //www gov.si/drzavni-organi/organi-v-sestavi/urad-za-kemikalije/
Controls on placing on the market of treated articles	Chemicals Office of the Republic of Slovenia	https //www gov.si/drzavni-organi/organi-v-sestavi/urad-za-kemikalije/

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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants	50	36	56	35	55	33	59	38	48	26	60	35	69	33
MG 2 Preservatives	6	4	7	3	6	3	3	1	3	1	3	1	3	1
MG 3 Pest control	41	30	45	28	49	31	55	25	45	33	58	24	58	35
MG 4 Other biocidal products	6	5	3	1	2	1	3	1	2	/	4	1	4	1

- **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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants	45	34	53	35	54	33	56	38	48	26	48	35	59	33
MG 2 Preservatives	5	4	5	3	5	3	5	1	3	1	3	1	3	1
MG 3 Pest control	38	28	37	23	36	31	40	25	35	23	58	24	55	35
MG 4 Other biocidal products	4	3	4	1	4	1	4	1	2	/	4	1	4	1

- **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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants	45	34	53	35	54	30	56	30	48	19	48	25	59	24
MG 2 Preservatives	5	4	5	3	5	2	5	1	3	1	3	1	3	1
MG 3 Pest control	38	28	37	23	36	15	40	19	35	22	58	19	55	31
MG 4 Other biocidal products	4	3	4	1	4	1	4	1	2	/	4	1	4	1

- **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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants	10	9	12	11	13	13	14	11	9	8	13	11	14	10
MG 2 Preservatives	1	1	1	1	/	/	/	/	1	1	1	1	/	/
MG 3 Pest control	9	8	7	7	7	6	8	6	9	7	12	10	12	9
MG 4 Other biocidal products	/	/	/		1	1	/	/	/	/	/	/	1	/

- **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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants	/	/	/	/	/	/	/	/	/	/	63	33	92	48
MG 2 Preservatives	/	/	/	/	/	/	/	/	/	/	/	/	/	/
MG 3 Pest control	/	/	/	/	/	/	/	/	/	/	/	/	/	/
MG 4 Other biocidal products	/	/	/	/	/	/	/	/	/	//	/	/	/	/

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)

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All inspections of biocidal products carried out by Chemicals Office of the Republic of Slovenia include the control of status of biocidal product in the national Register of biocidal products. National Register is being updated regularly (monthly) and contains all  biocidal products (BPs) authorised by CA regardless the procedure; consequently, the Chemicals  inspection can not estimate  the proportion of the BPs regarding the process resulting the registration into national Register.
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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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants														
MG 2 Preservatives														
MG 3 Pest control														
MG 4 Other biocidal products														

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

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Inspections od formulators of BP cover the control of Letter of Access.
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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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants														
MG 2 Preservatives														
MG 3 Pest control														
MG 4 Other biocidal products														

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

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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants	50	36	56	35	55	33	59	38	48	26	60	35	69	33
MG 2 Preservatives	6	4	7	3	6	3	3	1	3	1	3	1	3	1
MG 3 Pest control	41	30	45	28	49	31	55	25	45	33	58	24	58	35
MG 4 Other biocidal products	6	5	3	1	2	1	3	1	2	/	4	1	4	1

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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants			1	/			2	/			1	/	2	/
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

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were not carried out
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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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 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)**

```
were not carried out
```

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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 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)

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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants	10	7	/	/	2	1	3	2	2	1	10	4	2	1
MG 2 Preservatives														
MG 3 Pest control														
MG 4 Other biocidal products														

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

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 2013	NC 2013	Total 2014	NC 2014	Total 2015	NC 2015	Total 2016	NC 2016	Total 2017	NC 2017	Total 2018	NC 2018	Total 2019	NC 2019
MG 1 Disinfectants	10	7	/	/	2	1	3	2	2	2	10	4	2	1
MG 2 Preservatives														
MG 3 Pest control														
MG 4 Other biocidal products														

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)

In the electronic version of the Register of Poisonings with Biocidal Products of the Republic of Slovenia, from 2013 to 2019, 10.210 poisonings or selected poisons were reported (4430 reports of poisonings and 5785 calls to the 24-hour toxicology service of the Poison Control Center).

Table 1: Number of poisonings from 2013-2019

Year	No. of poisonings with biocidal products
2013*	43
2014	28
2015	22
2016	21
2017	21
2018	29
2019	48

Table 2: Age of persons exposed and poisoned to biocidal products (data are missing for 16 persons)

Age groups	No. of cases
00-02	54
02-18	61
18-30	19
30-50	43
50-70	19

Table 3: Overview of poisoning incidents from 2013-2019

2013* (n=1525)
2014 (n=1419)
2015 (n=1117)
2016 (n=1315)
2017 (n=1208)
2018 (n=1511)
2019 (n=2120)
See the table in the survey.

Table 4: Active substances in biocidal products to which persons have been exposed to or poisoned by (2013-2019) (numbers represent poisoning cases)

PT1	Ethanol	9
	Ethanol; biphenyl-2-ol	1
	Ethanol; propan-1-ol	6
	Ethanol; propan-2-ol	1
	Ethanol 96%	1
	Ethanol 96% ; 1-propanol	1
	Quaternary ammonium compounds, benzyl-C12-16-alkyldimethyl chlorides	2
	No data (active substance was not identified)	20
	Propan-1-ol ; propan-2-ol	1
	Propan-2-ol	1
	Triclosan	2
PT4		
	Didecyldimethylammonium chloride ; borate	1
	Didecyldimethylammonium chloride; propan-2-ol	1
	Quaternary ammonium compounds, benzyl-C12-16-alkyldimethyl chlorides, didecylammonium chloride glutarate; didecylammonium chloride; glutaraldehyde	1
	Sodium dichloroisocyanurate dihydrate	1
	Sodium hypochlorite	2
	No data (active substance was not identified)	4
	Hydrogen peroxide	2
PT18		
	Abamectin	2
	Alpha cypermethrin	2
	Bifentrin	1
	d-phenotryn ; tetramethrin	1
	Deltamethrin ; tetramethrin ; piperonyl butoxide	3
	Esbiotrin	1
	Geraniol	1
	Permetrhin	7
	Permethrin; tetramethrin ; piperonyl butoxide	1
	Prallethrin	4
	No information	8
	Spinosad	1
	Tetramethrin	1
	Transfluthrin	1
PT8		
	Boric acid; cooper(II)carbonate and cooper(II) hydroxide (1:1); alkil C12-16 dimetil benzil amonijev klorid	1
	Boric acid; cooper(II)carbonate and cooper(II) hydroxide (1:1); quaternary ammonium compounds, benzyl-C12-16-alkyldimethyl chloride	1
	Quaternary ammonium compounds, benzyl-C12-16-alkyldimethyl chlorides	3
	Permethrin ; 3-iodo-2-propinyl butylcarbamate	1
	Permethrin; propiconazole; 3-iodo-2-propinyl butylcarbamate	1

```
        No data (active substance was not identified)   5
PT15 No data (active substance was not identified)      1
PT2
        Didecyldimethylammonium chloride        2
        Didecyldimethylammonium chloride propan-2-ol    1
        Chlorine        14
        Quaternary ammonium compounds, benzyl-C12-16-alkyldimethyl chlorides    3
        Quaternary ammonium compounds, benzyl-C12-16-alkyldimethyl chlorides; hydrogen peroxide 4
        Sodium hypochlorite     12
        Polymer quaternary ammonium chloride    1
        No data 8
        Symclosene; cooper sulphate     1
        Tetraacetylene; disodium carbonate, hydrogen peroxide   1
        Hydrogen peroxide; quaternary ammonium compounds, benzyl-C12-16-alkyldimethyl chl       1
PT19    ethyl N-acethyl-N-buthyl-beta-alaninate 1
        Geraniol        1
        Icaridine       1
        N,N-diethyl-m-toluamide 1
        Pirethrins and pirethroids; ethanol     1
        No data (active substance was not identified))  7
PT14
        Brodifakum      1
        Bromadialone    1
        Bromadiolone    21
        No data (active substance was not identified)   11
PT5
        Dichloroizocianuric acid, Sodium salt (NaDCC)   1
        Chlorine        2
PT3
        Phoxim  2
        Acetic acid 80% 1
        No data (active substance was not identified)   1
PT10              No data (active substance was not identified)        2
PT not known  No data (active substance was not identified)     4
PT21               Cooper thiocyanate; zinc pyrithione 1


Severity of poisoning by biocidal products or exposure to biocidal products (2013-2019)
Severity of poisoning                                   Number of patients
Fatal intoxication                                              1 (ethanol + disinfectant; abuse)
Severe intoxication                                             1 (bromadiolone, suicide attempt)
Moderate poisoning                                               9
Mild poisoning                                                  67
Exposure without signs of intoxication        91
Assessment of severity was not possible      43

*Information for the year 2013 covers the whole year.
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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	0	22	0	13	0	13	0	10	1	8	1	14	0	30
MG 2 Preservatives	0	2	0	5	0	4	0	1	0	0	0	1	0	2
MG 3 Pest control	0	19	1	8	0	4	0	8	0	12	0	13	0	16
MG 4 Other biocidal products	0	0	0	1	0	1	0	2	0	1	0	0	0	0

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	2	5	5	8	7	5	4
Number of enquiries on biocidal products	155	485	476	796	721	534	408
Number of enquiries on treated articles	3	10	9	16	22	11	8
Total number of enquiries per year	160	500	490	820	750	550	420

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 advice to SMEs?

- (●) Yes
- ( ) No

Please indicate how the advice is customized for the needs of SMEs and refer to the corresponding links of the relevant websites for information

> The majority of companies in Slovenia are SMEs. Chemicals Office (CORS) supports all companies through Helpnet. As not all companies are members of Slovenian Chamber of Commerce, CORS organizes additional events free of charge, such as bilateral meetings, videoconferences, once per year a Chemical Safety day.

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 report.

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

Are certification procedures or training schemes in place (organised by e.g. eCAs, public authorities, sector organisations) for professional users of biocidal products?

- (●) Yes
- ( ) No
- ( ) 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	PT2 pools	National Institute for Public Health	once per year
2	PT5 drinking water	National Institute for Public Health	once per year
3	PT14 rodenticides PT18 insecticides	National Institute for Public Health	every 5 years

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 minimising their use? (Article 17(5) of the BPR)

- (●) Yes
- ( ) No

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	press conference	2015	presentation of the proper use of insecticides and repellents
2	different lectures	2017, 2018	safe use of BPs in pools, drinking water and for disinfection
3	implementing regulation	from 2018 onwards	restriction of professional use
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
- ( ) 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
1	flyer	2019	schools	information on excessive use of repelents
2	leaflet	2013 onwards	schools and others	classification and labelling of hazardous chemicals
3				
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	/				
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

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Data on poisoning incidents were revised in October 2020.
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Contact

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