

No	Service code	Service	Unit	Price without VAT, EUR	VAT, EUR	Price with VAT, EUR
1.1.	07-01.	Assessment of application to receive biocidal product inventory number	1 biocidal product	60,20	0,00	60,20
	07-01.01.	Assessment of application to receive biocidal product inventory number – additional fee for every biocidal product within biocidal product family	1 biocidal product	32,55	6,84	39,39
	07-11.	Assesment of detergent evaluation	1 mixture	12354,76	2594,50	14949,26
	07-13.	Assessment of authorisation and documentation of the identical biocidal product issued by Another EU Member State or European Economic Area competent authority for mutual recognition within one company	1 mixture	976,47	205,06	1181,53
	07-14.	Assessment of compliance with the definition of the biocidal product in accordance with Regulation (EU) No 528/2012	1 mixture	113,92	23,92	137,84
	07-16.	Confirmation of the issued inventory number	1 inventory number	55,33	11,62	66,95
	07-37.	Minor changes within the scope of the inventory provided (inventory number, product name, type of product if the active substance or concentration does not change)	1 biocidal product	65,10	13,67	78,77
	07-38.	Assessment of application to receive biocidal product inventory number in three work days	1 biocidal product	146,47	30,76	177,23

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Evaluation of application for approval of an active substance in accordance with Article 7 and 8 of Regulation (EU) No 528/2012						
1.2.	07-18.	Evaluation of application for approval of an active substance	1 active substance	122872,00	0,00	122872,00
1.3.	07-19.	Evaluation of application for approval of an active substance for an additional product type	1 product type	38931,14	0,00	38931,14
1.4. Evaluation of application for renewal of an active substance in accordance with Article 14 of Regulation (EU) No 528/2012						
1.4.1.	07-20.01.	Full evaluation of the application	1 active substance	122872,00	0,00	122872,00
1.4.2.	07-20.02.	Full evaluation of the application is not necessary	1 active substance	15136,97	0,00	15136,97
1.5.	07-21.	Evaluation of application for the authorisation of the same biocidal product in accordance with Article 17(7) of Regulation (EU) No 528/2012	1 biocidal product	3440,22	0,00	3440,22
1.6.	07-22.	Evaluation of application for the authorisation of the same biocidal product through mutual recognition, if related reference product is authorised in Latvia	1 biocidal product	963,26	0,00	963,26
1.7.	07-23.	Evaluation of application for comparative assessment of biocidal products in accordance with Article 23 of Regulation (EU) No 528/2012	1 biocidal product	17773,94	0,00	17773,94
1.8. Evaluation of application for authorisation made under a simplified authorisation procedure in accordance with Article 26 of Regulation (EU) No 528/2012						
1.8.1.	07-24.01.	Evaluation of application for single biocidal product	1 biocidal product	12026,34	0,00	12026,34
1.8.2.	07-24.02.	Evaluation of application for biocidal product family	1 biocidal product family	13750,48	0,00	13750,48
1.8.3.	07-24.03.	Evaluation of application for single same biocidal product	1 biocidal product	1115,62	0,00	1115,62

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1.9.	07-25.	Revision of labelling of biocidal product authorised in accordance with the simplified authorisation procedure (Article 27(1) of Regulation (EU) No 528/2012)	1 biocidal product	137,61	0,00	137,61
1.10. Evaluation of application for authorisation of biocidal product in accordance with the Article 29, 30 and 55 of Regulation (EU) No 528/2012						
1.10.1.	07-26.01.	Evaluation of application for single biocidal product	1 biocidal product	59216,60	0,00	59216,60
1.10.2.	07-26.02.	Evaluation of application for biocidal product family with 5 biocidal products	1 biocidal product family (5 biocidal products)	77048,15	0,00	77048,15
1.10.3	07-26.03.	Evaluation of application for biocidal product family for every next 5 biocidal products within 1 biocidal product family	Every next 5 biocidal products	278,91	0,00	278,91
1.10.4.	07-26.04.	Evaluation of the application for provisional authorisation in accordance with the Article 55 of Regulation (EU) No 528/2012	1 biocidal product	62390,94	0,00	62390,94
1.10.5.	07-26.05.	Evaluation of the application for authorisation of a restricted and controlled use of a biocidal product in specific cases in accordance with the Article 55 of Regulation (EU) No 528/2012	1 biocidal product	709,94	0,00	709,94
1.11	07-27.	Evaluation of application for authorisation of biocidal product or biocidal product family for additional product-type	1 product type	20554,92	0,00	20554,92

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1.12.	07-28.	Evaluation of application for additional biocidal product within a biocidal product family authorised	1 biocidal product	412,83	0,00	412,83
1.13. Evaluation of application for renewal of a national authorisation in accordance with Article 31 of Regulation (EU) No 528/2012						
1.13.1.	07-29.01.	Full evaluation of the application	1 biocidal product	59216,60	0,00	59216,60
1.13.2.	07-29.02.	Full evaluation of the application is not necessary	1 biocidal product	12109,57	0,00	12109,57
1.14.	07-30.	Evaluation of application for an authorisation of biocidal product through mutual recognition in sequence in accordance with Article 33 of Regulation (EU) No 528/2012	1 biocidal product/ 1 biocidal product family	2752,18	0,00	2752,18
1.15.	07-31.	Evaluation of application for an authorisation of biocidal product through mutual recognition in parallel in accordance with Article 34 of Regulation (EU) No 528/2012	1 biocidal product/ 1 biocidal product family	2752,18	0,00	2752,18
1.16. Evaluation of application for amendment of an authorisation						
1.16.1	07-32.01.	Evaluation of application in accordance with Article 50 of Regulation (EU) No 528/2012, if there is an administrative change	1 biocidal product	1394,53	0,00	1394,53
1.16.2. Evaluation of application in accordance with Article 50 of Regulation (EU) No 528/2012, if there is a minor change						
1.16.2.1.	07-34.01.	Evaluation of application, if there is a minor change (the reference Member State)	1 biocidal product	3270,80	0,00	3270,80
1.16.2.2.	07-34.02.	Evaluation of application, if there is a minor change (the Member States concerned)	1 biocidal product	1039,56	0,00	1039,56

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1.16.3. Evaluation of application in accordance with Article 50 of Regulation (EU) No 528/2012, if there is a major change						
1.16.3.1.	07-35.01.	Evaluation of application, if there is a major change (the reference Member State)	1 biocidal product	11628,74	0,00	11628,74
1.16.3.2.	07-35.02.	Evaluation of application, if there is a major change (the Member States concerned)	1 biocidal product	1039,56	0,00	1039,56
1.16.3.3.	07-35.03.	Evaluation of application if there is a major change in authorisation issued in a simplified procedure	1 biocidal product	7208,89	0,00	7208,89
1.17	07-32.04.	Evaluation of application for parallel trade permit in accordance with Article 53 of Regulation (EU) No 528/2012	1 biocidal product	1376,09	0,00	1376,09
1.18. Evaluation of the dossier for the inclusion of the active substance in Annex I of Regulation (EU) No 528/2012 in accordance with Article 28 of Regulation (EU) No 528/2012						
1.18.1	07-36.01.	Evaluation of the dossier for the inclusion of the active substance in category 1, 2, 3, 4 or 5 of the Annex I of Regulation (EU) No 528/2012	1 chemical substance	35268,78	0,00	35268,78
1.18.2.	07-36.02.	Evaluation of the dossier for the inclusion of the active substance in category 6 of the Annex I of Regulation (EU) No 528/2012	1 chemical substance	93099,04	0,00	93099,04
1.19.	07-36.03.	Evaluation of application for the renewal of biocidal product having been subject to mutual recognition in accordance with Article 40 of Regulation (EU) No 528/2012	1 biocidal product	1673,43	0,00	1673,43