ANNEX I

Fees relating to active substances

Table 1

Standard fees

General description of task; relevant provision in Regulation (EU) No 528/2012	Specific condition or task description	Fee (EUR)
Approval of an active substance; Article 7(2)	Fee for the first product-type for which that active substance is approved	120.000
	Additional fee per additional product-type	40.000
	Additional fee per product-type (for both the first product-type and any additional product-type) if the active substance is a candidate for substitution in accordance with Article 10 of Regulation (EU) No 528/2012	20.000
	Fee for the amendment of an approval, other than the addition of a product-type.	20.000
Renewal of an approval; Article 13(3)	Fee for the first product-type for which renewal of that active substance is sought	15.000
	Additional fee per additional product-type	1.500
	Additional fee for the first product-type for which renewal of that active substance is sought in case a full evaluation is found necessary in accordance with Article 14(1) of Regulation (EU) No 528/2012	25.000
	Additional fee per additional product-type in case a full evaluation is found necessary in accordance with Article 14(1) of Regulation (EU) No 528/2012	2.500
	Additional fee per product-type (for both the first product-type and any additional product-type) if the active substance is a candidate for substitution in accordance with Article 10 of Regulation (EU) No 528/2012	20.000
Inclusion in Annex I of an active substance; Article 28	Fee for the first inclusion in Annex I of an active substance	10.000
	Fee for the amendment of an inclusion of an active substance in Annex I	2.000
Notification in accordance with Article 3a of Regulation (EC) No 1451/2007	Fee per substance/product-type combination. The fee for the notification shall be deducted from the subsequent application in accordance with Article 7 of Regulation (EU) No 528/2012.	10.000

Table 2

Fee reductions for applications for the approval, renewal of approval, inclusion in Annex I of active substances if the active substance manufacturer is an SME established in the Union, except where the active substance is a candidate for substitution

Type of enterprise	**Reduction (% of the standard fee)**
Micro enterprise	60
Small enterprise	40
Medium enterprise	20

ANNEX II

Fees for Union authorisation of biocidal products

Table 1

Standard fees

General description of task; relevant provision in Regulation (EU) No 528/2012)	**Specific condition or task description**	**Fee (EUR)**
Granting of Union authorisation, single product; Article 43(2)	Fee per product not identical with (one of) the representative product(s) assessed for the purpose of the substance approval	80.000
	Fee per product identical with (one of) the representative product(s) assessed for the purpose of the substance approval	40.000
	Additional fee per product when comparative assessment in accordance with Article 23 of Regulation (EU) No 528/2012 is required	40.000
	Additional fee per product when the requested authorisation is provisional in accordance with Article 55(2) of Regulation (EU) No 528/2012	10.000
Granting of Union authorisation, biocidal product family; Article 43(2)	Fee per family	150.000
	Additional fee per family when comparative assessment in accordance with Article 23 of Regulation (EU) No 528/2012 is required	60.000
	Additional fee per family when the requested authorisation is provisional in accordance with Article 55(2) of Regulation (EU) No 528/2012	15.000
Notification to the Agency of an additional product within a biocidal product family; Article 17(6)	Fee per additional product	2.000
Union authorisation of a same biocidal product;	Fee per product constituting a "same product" within the	2.000

Article 17(7)	meaning of [the Same Products Regulation]	
Major change of an authorised product or product family; Article 50(2)	Fee per application	40.000
Minor change of an authorised product or product family; Article 50(2)	Fee per application	15.000
Administrative change of an authorised product or product family; Article 50(2)	Fee per notification	2.000
Recommendation on the classification of a change of an authorised product or product family; Article 50(2)	Fee per request in accordance with [the Regulation on Changes]. If the recommendation is to classify the change as an administrative or minor change, the fee for the request shall be deducted from the subsequent application or notification in accordance with [the Regulation on Changes].	2.000
Renewal of Union authorisation, single product; Article 45(3)	Fee per product	5.000
	Additional fee per product in case a full evaluation is found necessary in accordance with Article 14(1) of Regulation (EU) No 528/2012	15.000
	Additional fee per product when comparative assessment in accordance with Article 23 of Regulation (EU) No 528/2012 is required	40.000
Renewal of Union authorisation, biocidal product family; Article 45(3)	Fee per product family	7.500
	Additional fee per product family in case a full evaluation is found necessary in accordance with Article 14(1) of Regulation (EU) No 528/2012	22.500
	Additional fee per product family when comparative assessment in accordance with Article 23 of Regulation (EU) No 528/2012 is required	60.000

Table 2

Fee reductions for applications for the granting and renewal of Union authorisation of biocidal products or biocidal product families, if the prospective authorisation holder is an SME established in the Union, except where the product contains an active substance which is a candidate for substitution

Type of enterprise	**Reduction (% of the standard fee)**
Micro enterprise	30
Small enterprise	20
Medium enterprise	10

ANNEX III

Other fees

General description of task; relevant provision in Regulation (EU) No 528/2012)	**Specific condition or task description**	**Fee (EUR)**
Technical equivalence; Article 54(3)	Fee, when difference between the active substance sources is limited to a change in manufacturing location, and application is based solely on analytical data	5.000
	Fee, when difference between the active substance sources goes beyond a change in the manufacturing location, and application is based solely on analytical data	20.000
	Fee when previous conditions are not met.	40.000
Annual fee for biocidal products authorised by the Union; Article 80(1)(a)	Fee per Union authorisation of a biocidal product	10.000
	Fee per Union authorisation of a biocidal product family	20.000
Mutual Recognition Submission fee; Article 80(1)(a)	Fee per product or product family concerned by an application for mutual recognition, per Member State where mutual recognition is sought	700
Appeal; Article 77(1)	Fee per appeal	2.500
Submission for inclusion in the list of relevant persons; Article 95	Fee per submission of a letter of access to a dossier already found complete by the Agency or an evaluating Competent Authority	2.000
	Fee per submission of a letter of access to part of a dossier already found complete by the Agency or an evaluating Competent Authority, together with complementary data	20.000
	Fee per submission of a new dossier	40.000
Requests under Article 66(4) submitted to the Agency	Fee per item for which confidentiality is requested	1.000