EUROPEAN
COMMISSION

Brussels, XXX
PLAN/2024/989 ANNEX
(POOL/E4/2024/989/989-EN
ANNEX.docx)
[…](2025) XXX draft

ANNEXES 1 to 3

ANNEX

to the

COMMISSION IMPLEMENTING REGULATION (EU) .../...

amending Implementing Regulation (EU) No 564/2013 as regards the adaptation of fees to inflation

ANNEX I

Annexes I and II of Regulation (EU) No 564/2013 are amended as follows:

(1) in Annex I, Table 1 is replaced with the following:

'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	143 400
	Additional fee per additional product-type	47 800
	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	23 900
	Fee for the amendment of an approval, other than the addition of a product-type.	23 900
Renewal of an approval; Article 13(3)	Fee for the first product-type for which renewal of that active substance is sought	17 925
	Additional fee per additional product-type	1 793
	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	29 875
	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	2 988
	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	23 900
Inclusion in Annex I of an	Fee for the first inclusion in Annex I of an active	11 950

active substance; Article 28	substance	
	Fee for the amendment of an inclusion of an active substance in Annex I	2 390
Notification in accordance with Article 17(4) of Regulation (EU) No 1062/2014	Fee per substance/product-type combination. The fee for the notification shall be deducted from the subsequent application for approval	11950

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(2) In Annex II, Table 1 is replaced with the following:

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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	95 600
	Fee per product identical with (one of) the representative product(s) assessed for the purpose of the substance approval	47 800
	Additional fee per product when comparative assessment in accordance with Article 23 of Regulation (EU) No 528/2012 is required	47 800
	Additional fee per product when the requested authorisation is provisional in accordance with Article 55(2) of Regulation (EU) No 528/2012	11 950
Granting of Union authorisation, biocidal product family; Article 43(2)	Fee per family	179 250
	Additional fee per family when comparative assessment in accordance with Article 23 of Regulation (EU) No 528/2012 is required	71 700
	Additional fee per family when the requested authorisation is provisional in accordance with Article 55(2) of Regulation (EU) No 528/2012	17 925
Notification to the Agency of an additional product within a biocidal product family; Article 17(6)	Fee per additional product	2 390
Union authorisation of a same biocidal product; Article 17(7)	Fee per product constituting a ‘same product’ within the meaning of Commission Implementing Regulation (EU) No 414/2013 of 6 May 2013 specifying a procedure for	2 390

	the authorisation of same biocidal products in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([1])	
Major change of an authorised product or product family; Article 50(2)	Fee per application	47 800
Minor change of an authorised product or product family; Article 50(2)	Fee per application	17 925
Administrative change of an authorised product or product family; Article 50(2)	Fee per notification	2 390
Recommendation on the classification of a change of an authorised product or product family; Article 50(2)	Fee per request in accordance with Regulation (EU) No 354/2013. 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 Regulation (EU) No 354/2013.	2 390
Renewal of Union authorisation, single product; Article 45(3)	Fee per product	5 975
	Additional fee per product in case a full evaluation is found necessary in accordance with Article 14(1) of Regulation (EU) No 528/2012	17 925
	Additional fee per product when comparative assessment in accordance with Article 23 of Regulation (EU) No 528/2012 is required	47 800
Renewal of Union authorisation, biocidal product family; Article 45(3)	Fee per product family	8 963
	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	26 888
	Additional fee per product family when comparative assessment in accordance with Article 23 of Regulation (EU) No 528/2012 is required	71 700

([1])

OJ L 125, 7.5.2013, p. 4., ELI: http://data.europa.eu/eli/reg/2012/528/oj

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ANNEX II

‘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	5 975
	Fee, when difference between the active substance sources goes beyond a change in the manufacturing location, and application is based solely on analytical data	23 900
	Fee when previous conditions are not met.	47 800
Annual fee for biocidal products authorised by the Union; Article 80(1)(a)	Fee per Union authorisation of a biocidal product	11 950
	Fee per Union authorisation of a biocidal product family	23 900
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	837
Appeal; Article 77(1)	Fee per appeal	2 988
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	2 390
	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	23 900
	Fee per submission of a new dossier	47 800
Requests under Article 66(4) submitted to the Agency	Fee per item for which confidentiality is requested	1 195

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ANNEX […]