

# Ctgb Tariffs Decree per 1 January 2026

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Decree of the Board for the Authorisation of Plant Protection Products and Biocidal Products published on 24 September 2025, to establish the tariffs (i.e. fees and charges), payable in connection with the performance of its statutory tasks and other services, as described in this Decree and the corresponding annexes: (Tariffs Decree 2026)

The Board for the Authorisation of Plant Protection Products and Biocidal Products,

- By virtue of Article 74 Regulation (EC) 1107/2009 and Article 80 Regulation (EU) No 528/2012, Article 10, first clause, *Wet gewasbeschermingsmiddelen en biociden* (Plant Protection Products and Biocidal Products Act), as well as Article 17 *Kaderwet zelfstandige bestuursorganen* (Independent Administrative Bodies Framework Act);
- Taking into consideration that the transitional provisions in Regulation (EC) 1107/2009 require that fees and charges must also continue to be set for processing applications for the authorisation of plant protection products that must be processed according to the provisions in the now expired Directive 91/414/EEC and its implementation in the *Wet gewasbeschermingsmiddelen en biociden* (Plant Protection Products and Biocidal Products Act), as applicable before 14 June 2011;
- Taking into consideration that the implementation of Regulation (EU) 528/2012 resulted in additional arrangements regarding the tariffs (fees and charges), invoicing and payment for the services and activities that the Ctgb provides under the Regulation;
- Taking into consideration that the transitional provisions in Regulation (EU) 528/2012 require that fees and charges must also continue to be set for processing applications for the authorisation of biocidal products that must be processed according to the provisions in the now expired Directive 98/8/EC or on the basis of the *Wet gewasbeschermingsmiddelen en biociden* (Plant Protection Products and Biocidal Products Act) as applicable before the entry into force of the Act of 6 November 2013 to amend the Plant Protection Products and Biocidal Products Act with the aim of implementing the Biocidal Products Regulation;

Decree:

## Chapter 1. General

### Article 1. Terms and definitions

For the purpose of this Decree, the following definitions shall apply:

- a. Plant Protection Products Regulation: Regulation (EC) 1107/2009 of the European Parliament and the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC
- b. Plant Protection Directive: Council Directive 91/414/EEC of 15 July 1991 concerning the placing of plant protection products on the market.
- c. Biocidal Products Regulation: Regulation (EU) 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the placing of biocidal products on the market and the use of biocidal products
- d. Biocidal Products Directive: Directive 98/8/EC of the European Parliament and the Council of 16 February 1998 concerning the placing of biocidal products on the market
- e. the Act: Plant Protection Products and Biocidal Products Act (Wgb)
- f. the Regulation: Regulations on plant protection products and biocidal products (Rgb)
- g. Active substance:
  - i. plant protection products
    1. an active substance as referred to in Article 30 of the Plant Protection Products Regulation, or
    2. an active substance which
      - a. according to Implementing Regulation (EU) 540/2011 was not approved and
      - b. was not yet on the market on 15 July 1993 and
      - c. was therefore not made equivalent pursuant to a community measure
  - ii. biocidal products
    1. active substance as referred to in Article 3.1.e of the Biocidal Products Regulation
- h. Register for Biocidal Products (R4BP): Information system for submitting applications for the authorisation of biocidal products under Regulation (EU) 528/2012 and for exchanging information between authorities, the Commission, Agencies and applicants.
- i. Service Desk (SD) consists of a front and back office: front office concerns the first-line support provided by the Service Desk, while the back office concerns the second-line support provided by various expertises.
- j. Actual costs incurred: they are the internal costs incurred by the Ctgb (the established hourly rate, multiplied by the number of hours spent by the Ctgb) and the actual costs of third parties (including the VAT paid to such third parties). More information about the working method and cooperating partners can be found on the website [www.ctgb.nl](http://www.ctgb.nl).
- k. Timely payment means that the payment is received by the Ctgb before the payment deadline specified on the invoice.

## Chapter 2. Fees and charges

### Article 2. Fees and charges

The Ctgb levies fees and other charges in connection with the performance of its statutory tasks and other services. The hourly rate for 2026 has been set at €202. The Service Desk fees are listed in Annex I, the application fees for plant protection products are listed in Annex II, and those for biocidal products are listed in Annex III.

### Article 3. Fixed fees and charges, and fees and charges based on subsequent costing

1. Due to the differences between the various types of applications, the Ctgb applies two types of fees and charges:
  - a. Applications based on **subsequent costing**. An **advance payment** is requested for this. The final settlement takes place at the end of the application process based on actual costs incurred. If the advance payment is insufficient, additional charges will be invoiced. If applications for plant protection products are withdrawn before the dossier is formally submitted, the start-up costs already paid will not be refunded.
  - b. Applications based on a fixed fee. These fees and charges must be paid in advance and no final settlement will be made. If the Ctgb decides to request additional information, the costs for assessing this additional information will be charged. Regarding biocidal product applications that the Ctgb processes as CMS (Concerned Member State), several years may have passed between the submission of the application and the start of the evaluation. This is the time needed by the evaluating Member State to complete the initial assessment. The fees and charges that are invoiced by the Ctgb as CMS for these applications will be based on the costs as indicated in the Ctgb Tariffs Decree for the year in which the evaluation starts at Ctgb. This applies to the entire amount, i.e. start-up costs as well as evaluation costs. Previously paid start-up costs will be deducted.
2. For work for which no fees and charges have been set in advance, the actual costs will be charged.

### Article 4. Annual fee

1. For the registration of an authorised plant protection product or biocidal product, an annual fee is charged.
2. The reference date for the registration of an authorised plant protection product or biocidal product is 1 February, which is also the date on which the annual registration fee for the following year becomes due if the registration is not withdrawn in writing before this date.
3. If the applicant does not pay the annual fee or does not pay in full before this date, this applicant is legally in default according to Article 6:81 of the Netherlands Civil Code.

#### **Article 5. Frame formulation**

Regarding applications for biocidal products under transitional law, it is possible to apply for authorisations for a group of very similar products by submitting a combined application.

### **Chapter 3. Invoicing, payment and subsequent costing/refund**

#### **Article 6. - Plant protection products and Biocidal products under transitional legislation in the Netherlands**

1. The fee payable to the Ctgb is calculated based on the specified items and tariffs.
2. To cover the start-up and assessment costs that are due, the Ctgb will send an invoice with a payment term of 30 days. Late payment may lead to declaring the application non-admissible.
3. If an advance payment has been made for the actual costs incurred, this payment is subtracted from the amount due. If the actual costs incurred are less than the advance payment, the difference is refunded to the applicant within 30 days after date of the final invoice. If the actual costs are higher than the advance paid, an additional invoice follows with a payment term of 30 days.
4. For other fees or charges, the applicant shall pay the invoice within 30 days of the invoice date, indicating the invoice number.

#### **Article 7. - Biocidal Products Regulation**

1. After receiving the application, the Ctgb calculates the fee that is payable on the basis of the specified items and tariffs. The invoice is sent as soon as possible following receipt of the application in R4BP.
  2. The Biocidal Products Regulation provides that the costs of an application procedure must be paid within 30 days after the invoice date. Additional time for payment cannot be provided.
  3. If payment is late, the application is rejected.
  4. If an advance payment has been made for the actual costs incurred, this payment is subtracted from the amount due. If the actual costs incurred are less than the advance payment, the difference is refunded to the applicant within 30 days after date of the final invoice. If the actual costs are higher than the advance paid, an additional invoice follows with a payment term of 30 days.
  5. If multiple applications are submitted simultaneously, for which a cluster assessment involving one or more expertises can be made for multiple applications, at the request of the applicant(s) the Ctgb can charge a reduced rate for these expertises (in accordance with Article 80.3.d).
- If there are multiple applicants, each applicant is jointly and severally liable for paying the entire amount to the Ctgb. The Ctgb does not become involved with the mutual payment arrangements between applicants.

## Chapter 4 – Other provisions

### **Article 8. Interim amendment of the Ctgb Tariffs Decree**

The Tariff Decree may, if necessary, be amended in the interim.

### **Article 9. Transitional regime**

Applications that were pending on the date this Decree went into force are processed from that date according to the provisions in this Decree.

### **Article 10. Withdrawal of previous Tariffs Decree**

The Ctgb Tariffs Decree 2025 shall be withdrawn.

### **Article 11. Entry into force**

This Decree shall take effect, retroactively when necessary, on 1 January 2026, following approval by Our Minister and publication of this Decree in the Staatscourant (Government Gazette).

### **Article 12. Official title**

The official title of this Decree is: “**Ctgb Tariffs Decree 2026**”.

Approved by the Board for the Authorisation of Plant Protection Products and Biocidal Products  
**date 24 September 2025**

The Board for the Authorisation of Plant Protection Products and Biocidal Products,  
represented by,

the Chair

## Explanation of the Tariffs Decree

### General

The Act stipulates that the costs incurred by the Ctgb during the implementation of its activities must be covered by the tariffs charged, which are to be set by the board. The fees and charges in this Decree are related directly to the costs incurred by the Ctgb during this implementation and require the approval of the Minister of Agriculture, Fisheries, Food Security and Nature (LNV).

### How the fees and charges are determined

The fees and charges cover the internal costs of the organisation. Applicants pay a fee that is based on the various phases and expertises that are relevant to the assessment of their application. Examples of phases are: intake/validation, assessment and additional questions. Expertises include, but are not limited to: project management, human toxicology, residues, ecotoxicology, environmental behaviour and environmental fate, chemistry and efficacy. Due to the diversity in magnitude and complexity of an application dossier, the costs can vary widely, depending on the amount of work required. The type of application also determines the costs.

The tariffs are established by this Decree. If the application fees and charges are invoiced based on advance payment and subsequent costing, the amounts mentioned are averages. The fees and charges for a specific application could therefore be higher or lower than this average. For further information, go to: [www.ctgb.nl](http://www.ctgb.nl).

### Payment in instalments

It is possible to pay in instalments. For more information, see the Ctgb website [www.ctgb.nl](http://www.ctgb.nl) or contact the Finance & Control department ([finance@ctgb.nl](mailto:finance@ctgb.nl)).

### Collection options

It is possible to have invoice payments automatically transferred to the Ctgb. For more information, see the Ctgb website [www.ctgb.nl](http://www.ctgb.nl) or contact the Finance & Control department ([finance@ctgb.nl](mailto:finance@ctgb.nl)).

### Communication

From the moment an application for authorisation under the Biocidal Products Regulation is submitted, all correspondence on costs and payment is done in the Register for Biocidal Products (R4BP).

## Addendum I - Fees and charges, Service Desk

Requests	Description	Amount
Category		
1	Service Desk questions answered by the front office	No fee charged
2	Service Desk question answered by the back office	<p>€300</p> <p>If the actual costs incurred are higher, then the total costs will be invoiced.</p> <p>If the hourly expenditure is estimated to exceed 14 hours, an advance payment of €3,000 will be requested.</p>
3	Pre-submission meeting (PSM) Meeting for questions (or sub-questions) about dossier preparation	Actual costs incurred with an advance payment of €1,700 per expertise to be involved in the meeting
4	Workshop Meeting to keep applicants informed of current developments in practice.	Actual costs incurred are determined for each workshop

## Addendum II - Application fees and charges for plant protection products

Applications relating to active substances and plant protection products are based on subsequent costing with an advance payment that is invoiced at specific times. The **advance payments** as mentioned in the tables hereafter are indicative and are based on the average costs of assessing these dossiers in past years. At the start of the application process, the initial assessment costs are estimated based on the content of the application in question. The advance payment to be invoiced may therefore differ from the amounts listed below. Assessment costs are invoiced when new data can be submitted for assessment. This takes place at four different points in time.

### 1. Application for active substance or Maximum Residue Limit (MRL)

#### a. Application for active substance

Type of application	Description	Advance
TGEURAP TGEURAPR	<b>Application for approval, amendment or renewal of the approval of an active substance for plant protection with the Netherlands as rapporteur</b>	
	Start-up costs (performing the admissibility check)	€36,000
	Assessment costs (prepare DAR or RAR, peer review and European decision-making)	€468,000
TGEURAPM TGEURPRM	<b>Application for approval, amendment or renewal of the approval of an active substance for plant protection based on a micro-organism with the Netherlands as rapporteur</b>	
	Start-up costs (performing the admissibility check)	€44,000
	Assessment costs (prepare DAR or RAR, peer review and European decision-making)	€229,000
TGEUCORA	<b>Application for approval, amendment or renewal of the approval of an active substance for plant protection with the Netherlands as co-rapporteur</b>	
	Start-up costs (provide commentary on substance assessment RMS)	€17,500
	Assessment costs (prepare DAR or RAR) <i>(only if parts of the DAR/RAR must be prepared in consultation with the RMS)</i>	€59,000
	Peer review and European decision-making	€29,000
TGEURPCD	<b>Application for assessment of confirmatory data for active substances with the Netherlands as rapporteur</b>	
	Assessment of confirmatory data	€28,000
TGDM	<b>Application for data matching of an active substance for plant protection</b>	€13,000
TGDMcat4	<b>Application for data matching of category 4 data for an active substance for plant protection</b>	€2,400

#### b. Application for Maximum Residue Limit (MRL)

Type of application	Description	Advance
MRL	<b>Application for derivation of the Maximum Residue Limit</b>	
	Start-up costs	€7,900
	Assessment costs	€28,000

## 2. Application for product

Type of application	Description	Advance
ZTG ZWTG ZKWTG ZRG	<b>Applications for zonal authorisation, minor change or renewal of a plant protection product with the Netherlands as rapporteur</b>	
	Start-up costs	€31,000
	Assessment costs	€118,000
ZWTG-LR ZTG-LR	Start-up costs	€24,000
	Assessment costs	€61,000
NLWERGZ NLKUG NLWG	<b>National applications via mutual recognition, minor extended authorisations and changes</b>	
	Start-up costs	€13,000
	Assessment costs	€17,000
NLTG NLWTG NLRG	<b>Applications for zonal authorisation change or renewal of a plant protection product with the Netherlands as Concerned Member State</b>	
	Start-up costs	€18,500
	Assessment costs	€37,000
PWSG	<b>Application to change in production process of active substance and/or additional production location of the active substance. Equivalence not yet assessed by another Member State</b>	
	Start-up costs	€2,500
	Assessment costs	Actual costs incurred
MTR	<b>Application to derive standard under the Water Framework Directive</b>	
	Start-up costs	€2,500
	Assessment costs	Actual costs incurred

## 3. Other applications

This concerns all other applications at a fixed fee. The differences between these fees depend on the activities that are required. The costs shown in Table b can vary based on whether or not assessments are carried out.

### a. Other applications - fixed fee

Type of application	Description	Amount
WNAW	Reporting an administrative change for an authorised product - address details	No fee charged
INTR	Application for withdrawing the authorisation of a product	
EXV	Request for reduction for exporting an authorised or registered product	
VIG	Request for information about testing on vertebrate animals	€500
IKV	Reporting obligatory notification as specified in Regulation (EC) 1107/2009 Article 56, clause 4 or Regulation (EU) 528/2012 Article 47	
AG/VAG	Application for a new derived authorisation or the renewal thereof	€1,300
PAG/VPAG	Application for a new parallel trade permit or the renewal thereof	

## b. Other types of applications

Type of application	Description	Amount*
PMG	Add or change manufacturer and change manufacturing location of a plant protection product	No fee charged
WNT	Application to change the name of an authorised plant protection product	€500
WYG-AG	Application for administrative change for a derived authorisation or parallel trade permit for a plant protection product	
OT	Application to transfer one or more authorisations	
NLWATG	Application for administrative change to the legal instructions for use (language changes and restrictions on the scope of permitted use)	
PWSGC	Application to change in production process of active substance and/or additional production location of the active substance. Equivalence has been assessed by another Member State and published on CircaB	
PGN	Application for authorisation of a trial for a plant protection product where treated crops <u>do not</u> enter the food chain	
TT-W	Application to change an adjuvant in a plant protection product	
UAG	Application to expand the scope of permitted use of a derived authorisation for a plant protection product	€800
UPAG	Application to expand the scope of permitted use of a parallel trade permit for a plant protection product	
PGR	Application to authorise a trial of a plant protection product where treated crops <u>do</u> enter the food chain (maximum of 5 crops; surcharge of 200 euros for each additional set of 5 crops)	
WYG	Applications for administrative changes to an authorised plant protection product – packaging/shelf life/labelling/other change to original or reference product	
WSGNW	Applications for administrative change for an authorised plant protection product – minor change in the formulation	€1,300
TT	Application for authorisation of an adjuvant in a plant protection product	
ART 38	Request for exemption of a plant protection product under Article 38 of the Wgb	Actual costs incurred

\*If an assessment is required, the actual costs incurred will be charged.

## 4. Annual fee

Annual fee	Annual fee	Amount
	Annual fee for a plant protection product authorised by the Ctgb	€2,210

## Addendum III – Fees and charges for applications involving biocidal products

### 1. Application under regulation (EU) No 528/2012 (BPR)

Applications relating to active substances and biocides under the BPR, where the Netherlands acts as evaluating Competent Authority (eCA/RMS), are invoiced according to subsequent costing with an advance payment that is invoiced at various times (see Chapters 2 and 3 for more information). The **advance payments** as mentioned in the tables hereafter are indicative and are based on the average costs of assessing these dossiers in past years. At the start of the application process, the initial assessment costs are estimated based on the content of the application in question. The advance payment to be invoiced may therefore differ from the amounts listed below.

Applications relating to biocides under the BPR, where the Netherlands acts as the Concerned Member State (CMS), are processed according to fixed fees and charges. These fees and charges are in force for all phases necessary for the assessment of a dossier.

The following categories are distinguished:

- a. Application for active substance with the Netherlands as rapporteur – subsequent costing
- b. Application with the Netherlands as evaluating Competent Authority (eCA) - subsequent costing
- c. Application with the Netherlands as Concerned Member State (CMS) – fixed fee

#### a. Application for active substance with the Netherlands as rapporteur – subsequent costing

Type of application	Case type R4BP	Description	Advance	
EU-B	AS-APP, AS-RNL, AS-EVA, AN-APP	<b>Application for approval of a new active substance or renewal of the approval of an active substance</b>		
EU-B	AS-APP, AS-RNL, AS-EVA, AN-APP	Start-up costs	One product type (PT)	€22,500
			Additional fee per PT when approval for multiple PTs is applied for	€5,700
		Assessment costs	Dossier for an application for inclusion in Annexe I of Regulation (EU) 528/2012	€113,000
			Regular dossier	€227,000
Costs of EU decision-making process*		€82,000		

\* These are the costs for a single active substance-PT combination

#### b. Application for product with the Netherlands as evaluating Competent Authority (eCA) - subsequent costing

Type of application	Case type R4BP	Description	Advance payment single product	Advance payment biocidal family
B-UTN, B-UTH, B-UTFN, B-UTR, B-UTFR,	UA-APP	<b>Application for Union authorisation of a biocidal product or biocidal product family or re-registration of a biocidal product via a Union authorisation</b>		
		Start-up costs	€23,500	€29,000
		Assessment costs	€78,000	€147,000
		Costs of EU decision-making process	€31,500	€71,000
B-TN, B-TH, B-TFN,	NA-APP	<b>Application for national authorisation of a biocidal product or biocidal product family or re-registration via a national biocidal product authorisation</b>		
		Start-up costs	€23,500	€29,000
		Assessment costs	€66,000	€95,000
		Peer review and decision-making costs	€23,500	€35,000
B-ET, B-ETF	SA-APP	<b>Application for simplified authorisation of a biocidal product or biocidal product family</b>		
		Start-up costs	€3,700	€6,900
		Assessment costs	€21,000	€30,000
B-TR, B-TRF	NA-RNL	<b>Application for renewal of a nationally authorised biocidal product or biocidal product family</b>		
		Start-up costs	€11,500	€17,500
		Assessment costs	€29,000	€55,000
		Peer review and decision-making costs	€23,500	€35,000
B-GW, B-GWF	NA-MAC, SA-MAC,	<b>Application for a major change in the (Union) authorisation of a biocidal product or biocidal product family</b>		

B-UGW B-UGWF	UA-MAC	Start-up costs	€11,500	€17,500
		Assessment costs	€29,500	€41,000
		Peer review and decision-making costs / EU decision-making process costs/	€11,500	€17,500
B-KW, B-KWF	NA-MIC, SA-MIC	<b>Application for minor change in the authorisation of a biocidal product or biocidal product family</b>		
		Start-up costs, assessment costs, peer review and decision-making costs	€4,300	€5,300
B-FL	NA-NPF, SA-NPF	Notification of addition of one or more products to an authorised biocidal product family		€2,200

### c. Application for a product with the Netherlands as Concerned Member State (CMS) – fixed fee

Type of application	Case type R4BP	Description	Amount single product	Amount biocidal product family
B-TWENS B-TWEFNS	NA-MRS	<b>Application for national authorisation of a biocidal product or biocidal product family via sequential mutual recognition</b>		
		Costs (for start-up + evaluation)	€20,300	€24,300
		<i>Additional costs:</i>		
		Comparative assessment screening*	€3,200	€3,200
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SPC	n/a	€1,300
B-TWENP, B-TWEHP B-TWEFNP	NA-MRP	<b>Application for national authorisation of a biocidal product or biocidal product family or re-registration of a national authorisation of a biocidal product via parallel mutual recognition</b>		
		Start-up costs	€1,800	€1,800
		Evaluation costs	€18,500	€22,500
		<i>Additional costs:</i>		
		Comparative assessment screening*	€3,200	€3,200
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SPC	n/a	€1,300
B-TWER, B-TWEFR	NA-RNL	<b>Application for renewal of a nationally authorised biocidal product or biocidal product family</b>		
		Start-up costs	€1,800	€1,800
		Evaluation costs	€18,500	€22,500
		<i>Additional costs:</i>		
		Comparative assessment screening*	€3,200	€3,200
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SPC	n/a	€1,300
B-GWC, B-GWFC	NA-MAC	<b>Application for a major change in the authorisation of a biocidal product or biocidal product family</b>		
		Start-up costs	€1,800	€1,800
		Evaluation costs	€11,500	€17,000
B-KWC, B-KWFC	NA-MIC	<b>Application for minor change in the authorisation of a biocidal product or biocidal product family</b>		
		Start-up costs and evaluation costs	€2,400	€3,400

\* If Tier I and/or Tier II must be implemented after screening, the actual costs incurred will be charged.

## 2. Application under transitional law

If one or more active substances for the specified Product Types is not yet included on the Union list of approved substances or Annex I of Regulation (EU) No. 528/2012, then Transitional legislation is in force.

Fixed fees and charges will be invoiced for applications under Transitional legislation. Applications under transitional legislation for authorisation of biocidal products or renewal or major amendment of these authorisations can qualify for a limited assessment under specific conditions. The conditions are published on the Ctgb website.

For additional and/or other services and activities for which no fee has been set, the actual costs incurred will be invoiced.

### a. Application for full or limited assessment or application for renewal or minor change of an authorisation – fixed fee

Type of application	Description	Amount
TB, TBL, WB	<b>Application for a biocidal product authorisation or for a major change to such an authorisation under Dutch transitional law or by means of a limited assessment</b>	
	Start-up costs	€7,400
TB, WB	Assessment costs for a complete assessment	
	Efficacy	€2,200
	Physicochemical properties of the product and analytical methods	€1,300
	Add-on (additional charge) for multiple active substances	€1,600
	Criteria with respect to public health and operator	€2,200
	Environmental behaviour and environmental fate	€3,000
	Project supervision	€4,500
	Evaluation of summaries and studies concerning physicochemical properties, analytical methods, efficacy, human toxicology and environmental fate	Actual costs incurred
TBL, WB	Assessment costs for a limited assessment*	
	Efficacy	€900
	Physicochemical properties of the product and analytical methods	€500
	Criteria with respect to public health and operator	€1,200
	Environmental behaviour and environmental fate	€800
	Project supervision	€2,500
KB	<b>Application for establishing frame formulation</b>	
	Application for establishing frame formulation of biocidal products under Dutch transitional law	€4,000
TVB	Start-up costs	€1,800
	Assessment costs	Actual costs incurred
WBK	Application for minor change of a biocidal product authorisation under Dutch transitional law	€2,200

\* If limited assessment is possible; otherwise, the full assessment fee applies.

## 3. Other applications

This concerns all other applications at a fixed fee. The differences between these fees depend on the activities that are required. The costs shown in Table b can vary based on whether or not assessments are carried out.

### a. Other applications - fixed fee

Type of application	Case type R4BP	Description	Amount
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WNAW	NA-, SA-ADC	Reporting an administrative change for an authorised product - address details	No fee charged
INTR	NA-, SA-CCL	Application for withdrawing the authorisation of a product	
EXV		Request for reduction for exporting an authorised or registered product	€500
VIB	UE-, NE-, SE-NOT	Request for information about testing on vertebrate animals	
IKV-B		Reporting obligatory notification as specified in Regulation (EC) 1107/2009 Article 56, clause 4 or Regulation (EU) 528/2012 Article 47	
AB/VAB	PP-APP	Application for a new derived authorisation or the renewal thereof	€1,300
PAB/VPAB		Application for a new parallel trade permit or the renewal thereof	

#### b. Other types of applications

Type of application	Case type R4BP	Description	Amount*
WBA	n/a	Application for administrative change for an authorisation (or derived authorisation) of a biocidal product under Dutch transitional law	€500
B-AW B-AWF	NA-, SA-ADC, NA-, SA-TRS, NA-MRG	Application for administrative change for an authorised biocidal product (or biocidal product family) according to Regulation 354/2013	
B-EM	SN-NOT	Notification of a simplified authorisation of a biocidal product to place it on the market in the Netherlands	€800
PB	ET-NOT	Notification of experiment or trial	€1,300
B-ST/B-STF/B-STR/B-STRF	NA-, SA-BBS, NA-, SA-BBP, NA-RNL	Application for a new authorisation of a same biocidal product (or biocidal product family), or the renewal thereof, with the Netherlands as eCA	
ART 46		Request for exemption of a biocidal product based on Article 46 of the Wgb	Actual costs incurred

\*If an assessment is required, the actual costs incurred will be charged.

#### 4. Annual fee

Annual fee	Annual fee for a biocidal product authorised by the Ctgb	Amount
	Authorisation of a biocidal product	€1,700
	Authorisation biocidal family*	€1,700
	Additional charge per family member*	€200
	<i>*For placing biocidal products from a product family on the Dutch market, a fee will be charged for both the family and the individual members of the family.</i>	
	The maximum fee that is charged for a family per year.	€8,000

## Addendum IV - Most important changes

### General

Hourly rate 2026	The hourly rate for 2026 has been indexed by 3% and now totals €202.
Service Desk	A minimum fee of €300 is charged for a Service Desk question answered by the back office. If the actual costs incurred are higher, then the total costs will be invoiced.
other applications	With effect from 2026, a simpler system of fees and charges (with three categories) will be used, where the classification is based on the expected hourly expenditure.

### Plant protection

General	The fees and charges for applications for biocidal products have been adjusted as a result of the increase in the hourly rate indicated above.
Applications	The advance payments for the application types based on subsequent costing have been adjusted where necessary. These advance payments are based on the historical average cost of applications by type and the changes in activities due to new guidelines, among other considerations. In particular, the advance payments for EU substance assessments have been increased due to the greatly increased magnitude of these dossiers.
TGEURAP, TGEURAPM, TGEURAPR, TGEURAPRM	The distinction between the costs for summarising and reviewing (DAR/RAR), peer review and European decision-making has been discontinued. In this Tariffs Decree, all such activities are now classified as assessment costs. In addition, the fact that assessment fees are invoiced at four different points in time has been clarified.
Annual fee	The annual fee has been indexed by 9% and will be €2,210 in 2026. This indexation is due to three factors: a decrease in the number of authorised products, a periodic indexation of the hourly rate and an increase in the activities that are covered by the annual fee.

### Biocidal product

General	The fixed fees and charges for applications for biocidal products have been adjusted as a result of the increase in the hourly rate indicated above.
Applications	The advance payments for the application types based on subsequent costing have been adjusted where necessary. These advance payments are based on the historical average cost of applications by type and the changes in activities due to new guidelines, among other considerations.
Annual fee	The annual fee has been indexed by 9% and will be €1,700 in 2026. This indexation is due to three factors: a decrease in the number of authorised products, a periodic indexation of the hourly rate and an increase in the activities that are covered by the annual fee.