

# Rectification of the Ctgb Tariffs Decree 2020

## Content

General	
Chapter 1	Terms and definitions
Chapter 2	Other provisions
Chapter 3	Explanation of the Ctgb Tariffs Decree
Chapter 4	Invoicing, payment and subsequent costing/refund (Plant protection and Biocidal products under transitional law)
Chapter 5	Invoicing, payment and subsequent costing/refund Biocidal Products Regulation
Chapter 6	Most important changes relative to the Ctgb Tariffs Decree 2019
Chapter 7	Fees and charges, Service Desk
Chapter 8a	Fees and charges, Plant protection products (advance payment/subsequent costing) 1 Fees and charges, applications with active substance dossiers 2 Fees and charges, applications where Ctgb is zonal rapporteur Member State (zRMS) 3 Fees and charges, applications for national authorisation extended with minor uses (NLKUG)
Chapter 8b	Fees and charges, Plant protection products (fixed fees and charges) 1 Fees and charges, applications where Ctgb is Concerned Member State (CMS) 2 Fees and charges, mutual recognition of plant protection products (WERG) 3 National amendment application (NLWG) 4 Fees and charges, deriving maximal residue limit (MRL)
Chapter 8c	Fees and charges, Plant protection products, other applications 1 Fees and charges, other applications plant protection products, no assessment involved 2 Fees and charges, other applications plant protection products, for which additional assessment costs can be charged
Chapter 8d	Annual fee plant protection product
Chapter 9a	Fees and charges, Biocidal products (BPR) 1 Fees and charges, applications for approval of active substances 2 Fees and charges for applications for the authorisation of a product or for the amendment or renewal of a product authorisation, where the Ctgb is the evaluating competent authority (eCA) 3 Fees and charges for applications for the authorisation of a product or for the amendment or renewal of a product authorisation, where the Netherlands is a concerned Member State 4 Fees and charges for other or administrative requests
Chapter 9b	Fees and charges, Biocidal products (transitional law) 1 Fees and charges for product applications and other activities where one or more existing active substances are not yet included for the requested Product Types in the Union list of approved substances or Annex I of Regulation (EU) 528/2012 2 Other applications
Chapter 9c	Annual fee biocidal products

## General

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

- By virtue of Article 74 Regulation (EC) 1107/2009, 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 continue to apply to applications for the authorisation of plant protection products that must be processed according to the provisions in the now expired Directive (EEC) 91/414 and its implementation in the *Wet Gewasbeschermingsmiddelen en biociden* (Plant Protection Products and Biocidal Products Act), as in force before 14 June 2011;
- By virtue of Article 80 of Regulation (EU) 528/2012, Article 10, first clause, *Wet Gewasbeschermingsmiddelen en biociden* (Plant protection products and biocidal products Act), and Article 17 *Kaderwet zelfstandige bestuursorganen* (Independent Administrative Bodies Framework Act);
- Taking into consideration that the implementation of Regulation (EU) No. 528/2012 resulted in additional arrangements regarding the 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 also require that fees and charges continue to apply to applications for the authorisation biocidal products that must be processed according to the provisions in the now expired Directive (EC) 1998/98 or on the basis of the *Wet Gewasbeschermingsmiddelen en biociden* (Plant Protection Products and Biocidal Products Act) as in force before 6 November 2013 when it amended the Plant Protection Products and Biocidal Products Act for the implementation of the Biocidal Products Regulation;

**decree to establish the fees and charges for the performance of its statutory tasks and other services, as described in this decree and the corresponding annexes:**

## Chapter 1 - Definitions:

- 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
- 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
- f. the Regulation: Regulations on Plant Protection Products and Biocidal Products
- g. new active substance:
  - i. (plant protection products)
    - 1. an active substance as specified 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 on the market on 15 July 1993 and
      - c. was therefore not made equivalent pursuant to a Community measure
  - ii. (biocidal products) active substance as referred to in Article 3, first clause under e of the Biocidal Products Regulation
- h. zonal applications: applications for amendment/authorisation of plant protection products that are processed according to the zonal/interzonal assessment method described in Article 33 of the Plant Protection Products Regulation
- i. European applications: applications for approval or modification of the approval of an active substance, as referred to in Article 7 of the Plant Protection Products Regulation.

## Chapter 2 – Other provisions

### 1. Requests for public disclosure

Regarding requests for public disclosure, the Ctgb applies the provisions in the *Besluit tarieven openbaarheid van bestuur* (Decree fees and charges freedom of information) accordingly.

### 2. Transitional regime

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

### 3. Withdrawal of previous Tariffs Decrees

The “Ctgb Tariffs Decree 2019” is withdrawn.

### 4. Entry into force

This Decree shall take effect, retroactively to the extent necessary, on 1 January 2020, following approval by Our Minister and publication in the Staatscourant.

### 5. Interim amendment of the Ctgb Tariffs Decree

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

### 6. Official title

The official title of this Decree is “**Tarievenbesluit Ctgb 2020**” (Ctgb Tariffs Decree 2020).

Adopted by the Board for the authorisation of plant protection products and biocidal products dated 25 September 2019

The Board for the Authorisation of Plant Protection Products and Biocidal products,  
on his behalf,  
Chairman,

ir. J.F. de Leeuw

## Chapter 3 - Explanation of the Tariffs Decree

### General

The law stipulates that the costs incurred by the Ctgb in carrying out the assigned duties must be covered by the fees and charges to be set and charged by the Board. The fees and charges are directly related to the costs incurred by the Ctgb during implementation and require the approval of the Ministers for Agriculture, Nature and Food Quality (LNV) and Infrastructure and Water Management (IenW).

### Structure of fees and charges

The Ctgb works on the basis of the actual costs. These are the hours that experts spend assessing the dossier on safety for humans, animals and the environment. Applicants pay fees and charges consisting of the various assessments that are required for their applications. Due to the size and complexity of a dossier, such as use restricted to greenhouses or outdoors, whether it is an application for a single product or for a large biocidal product family, the costs can vary considerably.

The costs of an application consist of fees and charges for the various phases and aspects of the assessment, such as human toxicology, residues, ecotoxicology, behaviour and environmental fate, physicochemical properties and efficacy. The various phases are essentially stacked on top of each other. The costs depend on the required work, which differs per application.

The type of application also determines the costs. For example, a product that has already been authorised in another Member State can be placed on the market in the Netherlands by means of mutual recognition. The Ctgb will then use most of the assessment from the other EU Member State and will mainly look at aspects that are specific to the Netherlands. The costs for mutual recognition are considerably lower than for a complete assessment. However, if additional assessments are required for authorisation on the Dutch market, this can still increase the costs.

### Fixed fees and charges, and fees and charges based on advance payment and subsequent costing

Due to the differences between the various types of applications, the Ctgb applies two types of fees and charges: 1) fees and charges based on advance payment and subsequent costing, and 2) fees and charges based on fixed rates. Fees and charges must be paid in advance.

If the Board decides to request additional information, the costs for assessing this additional information will also be charged.

For work for which no fees and charges have been set in advance, the actual costs will be charged.

The fees and charges are established by the Tariffs Decree and relate to:

- *Applications for authorisation of an active substance of a plant protection product or for authorisation of an active substance according to (EU) 528/2012.*
- *Applications for authorisation (e.g. applications for new products, amendments or renewals of existing authorisations) of a biocidal product or plant protection product and other fees and charges (for example, annual fees, which authorisation holders must pay to the Ctgb based on Article 10, first paragraph of the Wgb).*
- *Requests for public disclosure by reference to the Tariffs Decree, Public Information Act (Besluit tarieven openbaarheid van bestuur).*
- *Service Desk requests.*
- *Exemption as referred to in Article 53 of the Plant Protection Products Regulation.*
- *Other activities.*

### Annual fee

For the registration of an authorised plant protection product and biocidal product, an annual fee is charged. The reference date for this annual fee is 1 February, which is also the date on which registration for the following year becomes payable if the registration is not withdrawn in advance in writing.

If the registrant does not pay the annual fee or does not pay it in full within the specified period, they are legally in default as stipulated in Article 6:81 of the Dutch Civil Code.

## Chapter 4 - Invoicing, payment and subsequent costing/refund (Plant protection products and Biocidal products under transitional law)

### Invoicing, payment and subsequent costing/refund

- a) The fee payable to the Ctgb is calculated based on the specified items and the corresponding fees and charges.
- b) The Ctgb will send an invoice for the application or assessment costs which is due within 30 days. Non-payment or late payment may lead to non-admissibility of the application under Article 2.3 of the Administrative Regulations 2018.
- c) 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.
- d) Notwithstanding the provisions in Article 4:87.1 of the *Algemene Wet Bestuursrecht (Awb)*, for fees and charges other than those referred to under b), the applicant must pay the invoice within 30 days after date of the invoice, stating the invoice number.

### Explanation of “Subsequent costing”

For various applications the costs are calculated based on subsequent costing of the actual costs incurred; if this is the case, this is expressly stated next to the corresponding application. For these applications, an advance payment will first be charged and then a final invoice will be made on the basis of subsequent costing and actual costs.

### Adjustment of the advance payment

Preceding the submission of an application, the prospective applicant can request the Ctgb to make an estimate of the actual costs of that application and adjust the advance payment accordingly; this estimate is based on substantiation provided by the prospective applicant. This advance payment can be changed if justified by the estimated hours.

### Explanation of “Actual costs”

The actual costs incurred 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 hired by the Ctgb as part of processing the application (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).

### Explanation of “Timely payment”

Timely payment means that the payment is received by the Ctgb before the payment deadline on the invoice or if it is shown that the amount to be paid has been transferred or deposited to the Ctgb before this deadline.

### Payment in instalments

Until two weeks after the invoice date, the applicant may request the Board for permission to pay the costs of the procedure at the Ctgb in instalments.

- This request will be granted if the costs of the procedure are more than €5,000.
- If the costs for the procedure are no more than €30,000, this amount may be paid in up to three monthly instalments of one-third of the total sum.
- If the costs for the procedure are more than €30,000, this amount may be paid in up to four monthly instalments of one-fourth of the total sum.

The first instalment must be paid within 30 days after the date of the original invoice. If payment is made in instalments, following timely payment of the first instalment, the Board formally accepts the application and notifies the applicant of the date of this acceptance.

### Collection options

It is possible to have invoice payments automatically transferred to the Ctgb. You can find more information about this on the website [www.ctgb.nl](http://www.ctgb.nl). For this, contact the Ctgb on [finance@ctgb.nl](mailto:finance@ctgb.nl).

### Frame formulation

The applicant for an authorisation under transitional law of a biocidal product can request the Ctgb to ascertain that the corresponding biocidal product qualifies as a frame formulation. The frame formulation offers the authorisation holder the possibility of obtaining a very similar authorisation (for example in different colours) more easily and at lower cost. The determination of the frame formulation entails additional application costs, which are added to the standard application costs.

## **Chapter 5 - Invoicing, payment and subsequent costing/refund Biocidal Products Regulation**

**With regard to biocidal products, the Ctgb must abide by the provisions regarding fees and charges laid down in the Biocidal Products Regulation as a matter of priority. These are explained below.**

### **Regulation (EU) 528/2012**

On 1 September 2013, Regulation (EU) No 528/2012 entered into force. This Regulation applies to all biocidal products and active substances in biocidal products (and applications for such products and substances) which, according to transitional legislation, do not fall under the operation of national law or Directive 98/8/EC. The Regulation contains rules about the amounts Member States may charge for the services they provide related to the procedures in the Regulation.

### **Basic assumptions and premises**

The Biocidal Products Regulation obligates Member States to take the following aspects into account when determining their fees and charges:

- the partial reimbursement of the fee if the applicant fails to submit the requested information within the deadline (Article 80 clause 3b);
- where appropriate, taking into account the specific needs of SMEs, including the possibility of splitting payments into several instalments and phases (Article 80 clause 3c);
- in the structure and amount of the fees, taking into account the circumstance of whether the information is submitted jointly or separately (Article 80 clause 3d);
- in duly justified cases, provided that the agency or the competent authority agrees, all or part of the fee may be waived (Article 80 clause 3e);
- under the Regulation, all communication between the applicant and the Ctgb takes place digitally;
- the Regulation introduces new types of applications and new types of authorisations, new obligations with respect to amounts charged, invoicing and payment, and the possibility of charging modified fees and charges;
- Member States that conduct the assessments must cover their costs;
- where the Biocidal Products Regulation does not offer a solution, Chapter 4 of this Ctgb Tariffs Decree applies.

### **Communication**

From the time an application is submitted, all correspondence on costs and payment takes place through the Register for Biocidal Products (R4BP).

### **Invoicing, payment and subsequent costing/refund (biocidal products)**

- After receiving the application, the Ctgb calculates the fee that is payable on the basis of the specified items, fees and charges. The invoice is placed in R4BP as soon as possible following receipt of the application.
- The Biocidal Products Regulation stipulates that the costs of an application procedure must be paid within a period of 30 days after the invoice date. Additional time for payment cannot be provided.
- If payment is late, the application is rejected.
- If the invoiced amount concerns an advance payment, the actual costs incurred are charged after the completion of the procedure by means of subsequent costing. If the actual costs incurred are less than the advance payment, the difference is refunded to the applicant within four weeks after date of the final invoice.

### **Reduced tariff for clustered applications**

If multiple applications are submitted simultaneously, for which a cluster assessment can be made on one or more aspects of multiple applications, at the request of the applicant the Ctgb can charge a reduced tariff for these aspects. 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.

### **Application of biocidal product families**

Under Regulation (EU) 528/2012 it is possible to apply for authorisation of several products at once. For applications where the Ctgb acts as evaluating Member State, an advance payment is charged. For mutual recognition of an authorisation of a biocidal product family by a different Member State, fixed fees and charges apply.

## Chapter 6 - Most important changes

### Crop protection

Comparative assessment	When the tariff decree was published, the tariffs were not yet known and the item listed here was for the record only. The rates have been set in accordance with this rectification.
Exemption as referred to in Article 53 of Regulation (EC) 1107/2009	The actual costs incurred will be charged. Advance payment will no longer be charged and graduated scales will no longer be used.

### Biocidal products

Application for renewal of a product authorisation (B-TWER)	The initial rate that was listed here was incorrect; this has now been amended.
Application for authorisation, extended authorisation, change/amendment and renewal (TB, UB, WSB, TVB)	Separate fees and charges were published for the assessment of criteria with respect to both public health and the operator. For the purpose of the assessment, this has been combined into one fee.

## Chapter 7 - Fees and charges Service Desk

Applications	Description	Amount
Category		
1	Up to 4 hours Service desk questions that can be processed within 4 hours.	No fee charged
2	4 hours and more Service desk questions that cannot be processed within 4 hours.	Actual costs incurred based on subsequent costing
3	Request for meeting (RFM) Meeting for potential applicants who want information about subsidiary questions concerning their dossier	Actual costs incurred based on subsequent costing
4	Short RFM Short meeting following a workshop, during which potential applicants can discuss the topic of the workshop in relation to their own dossier.	0.5 hours preparation + hours based on subsequent costing
5	Pre-submission meeting (PSM) During a PSM recommendations are made to the applicant regarding procedural and substantive aspects, based on a nearly complete dossier.	Actual costs incurred based on subsequent costing with an advance payment of €10,000
6	Briefing/informational meeting (general) Meeting with general information for potential applicants who do not yet have experience with an application and dossier.	€200
7	Workshop Meeting to keep applicants informed of current developments in practice.	For the record only

## Chapter 8a – Fees and charges Plant Protection Products (advance payment/subsequent costing)

### General

Regarding applications for active substances, zonal applications with the Netherlands as zonal Rapporteur Member State (zRMS) and NLKUG applications (National authorisation extended with minor uses), the fees and charges are based on **advance payment and subsequent costing**.

With these applications it is difficult to estimate in advance how many hours the assessment will take and there is a large spread in the required assessment time between the applications. The Ctgb invoices for the advance payment at the start of the application process.

1. Fees and charges, applications with active substance dossiers
2. Fees and charges, applications Ctgb as zonal Rapporteur Member State (zRMS)
3. Fees and charges, applications national authorisation extended with minor uses (NLKUG)

## 1. Fees and charges, applications with active substance dossiers

Type of application	Description	Amount
TGEURAP, TGEURAPR	<b>Ctgb rapporteur for application for initial approval or renewal of an active substance, excluding an active substance based on low-risk substances, micro-organisms or pheromones</b>	
	Application fee	€12,875
	Summarisation and assessment (preparation DAR or RAR)	Actual costs incurred with advance payment of €150,000*
	Activities related to the completion of an approval or renewal of an application for an active substance (Peer review; European decision-making)	Actual costs incurred with advance payment of €75,000*
TGEURAPM, TGEURPRM	<b>Ctgb rapporteur for application for initial approval or renewal of an active substance based on low-risk substances, micro-organisms, pheromones or a comparable active substance</b>	
	Application fee	€12,875
	Summarisation and assessment (preparation DAR or RAR)	Actual costs incurred with advance payment of €50,000*
	Activities related to the completion of an approval or renewal of an application for an active substance (Peer review; European decision-making)	Actual costs incurred with advance payment of €30,000*
TGEUCORA	<b>Ctgb Co-rapporteur substance dossier</b>	
	Fee for reviewing and commenting on substance assessment RMS	Actual costs incurred with advance payment of €20,000*
	Summarisation and assessment (preparation DAR or RAR) <i>(only if parts of the DAR/RAR must be prepared in consultation with the RMS)</i>	Actual costs incurred with advance payment of €50,000*
	Co-rapporteur activities to complete an approval or a renewal of the application for an active substance (peer review and European decision-making) <i>(only if parts of the DAR/RAR must be prepared in consultation with the RMS)</i>	Actual costs incurred with advance payment of €25,000*
TGEURPCD	<b>Ctgb rapporteur for the assessment of confirmatory data</b>	
	Assessment of confirmatory data	Actual costs incurred with advance payment of €15,500*
TGDM	Data matching for the active substance dossier Assessment of data matching for substance dossiers of non-notifiers of the EU authorisation of the active substance, including the determination of Category 4 data if applicable, as referred to in EU GD Art. 43 PPP renewals; SANCO/2010/13170 rev. 14 dated 7 October 2016.	Actual costs incurred with advance payment of €10,000*
TGDM-C4	Data matching of Cat. 4 data submitted later If Category 4 data have been claimed and determined in the primary assessment for data matching, then the applicant must provide these data before the set deadline and it must be assessed whether the data provided are sufficient to fully explain the data matching.	Actual costs incurred with advance payment of €2,000*

\* Advance payment is based on an average dossier. The advance payment can be changed if justified by the estimated hours. If the advance payment is insufficient to cover all costs, additional costs will be charged.

## 2. Fees and charges, applications Ctgb as zonal Rapporteur Member State (zRMS)

Type of application	Description	Amount
ZTG, ZTG-LR	<b>Zonal application for new authorisation NL = zRMS</b> The suffix –LR indicates a product application in which the low-risk substance procedure is followed.	Actual costs incurred with advance payment between €30,000 and €70,000.
ZWTG, ZWTG-LR	<b>Application to amend current authorisation</b> The suffix –LR indicates a product application in which the low-risk substance procedure is followed.	Actual costs incurred with advance payment between €30,000 and €70,000.
ZTHG	<b>Voluntary zonal renewal under transitional law.</b>	Actual costs incurred with advance payment between €30,000 and €70,000.
ZRG	<b>Renewal applications (Renewal of a product authorisation with the Netherlands as zRMS), for which postponement has been granted for submitting a dRR.</b>	
	<b>Application costs</b> This amount will be deducted from the advance payment after submitting the DRR.	€10,000
	Intake and assessment -Renewal application	Actual costs incurred with advance payment between €30,000 and €70,000.

\* Advance payment is based on an average dossier. The advance payment can be changed if justified by the estimated hours. If the advance payment is insufficient to cover all costs, additional costs will be charged.

### 3. Fees and charges, applications national authorisation extended with minor uses (NLKUG)

Type of application	Description	Amount
Applications national authorisation extended with minor uses (NLKUG)	Application to extend an existing authorisation in the Netherlands (not zonally) with minor uses only	Actual costs incurred with advance payment of €6,000*

\* Advance payment is based on an average dossier. The advance payment can be changed if justified by the estimated hours. If the advance payment is insufficient to cover all costs, additional costs will be charged.

## Chapter 8b – Fees and charges, Plant Protection Products (fixed fees and charges)

For the majority of assessments, costs are based on a **fixed fee**. This concerns applications for which the Netherlands acts as a Concerned Member State (CMS), mutual recognitions, NLWGs (applications for which only the national addendum is assessed) and MRL assessments. The first charge is for an application fee for the intake/commentary round. Next, it is assessed whether the dossier is valid and if no information is missing, after which the dossier can proceed to the assessment phase.

Applicants pay a fixed amount for the assessment based on the standard costs. 'Add-ons' (additional fees) are also charged for aspects such as verifying compliance with specific guidances, an additional draft Registration Report, the number of uses for the product, the plant-pest combination, assessment for an additional active substance or studies that must be evaluated.

1. Fees and charges, applications Ctgb as Concerned Member State (CMS)
  - 1.1. Application for new authorisation and renewal of an existing authorisation where the Netherlands acts as CMS (NLTG/NLRG)
  - 1.2. Application to amend current authorisation where the Netherlands acts as CMS (NLWTG)
2. Mutual recognition of plant protection products (WERG)
3. Application to amend current authorisation where only the national addendum needs to be assessed (NLWG)
4. Deriving maximum residue limit (MRL)

### 1. Fees and charges for applications where the Ctgb acts as Concerned Member State (CMS)

It is difficult to determine an average fee for this type of application, because different add-ons can be charged per application and the costs may therefore vary.

The assessment fees for the following types of applications are based on applications with 1 active substance, with up to 5 uses (GAP regulations) and 1 dRR (field or greenhouse use) per zone.

Additional fees are charged if a product contains multiple active substances, if there are more than 5 uses, if there is more than 1 dRR or when additional activities are required.

### 1.1 Application for new authorisation and renewal of an existing authorisation where NL acts as CMS (NLTG/NLRG)

An NLTG is an application for a new authorisation with the Netherlands as the Concerned Member State. An NLRG is a renewal of an authorisation dossier with the Netherlands as the Concerned Member State.

Type of application	Description	Amount
NLTG, NLRG	<b>Zonal application for new authorisation/renewal of authorisation where the Netherlands is CMS</b>	
	<b>Application fee for new authorisation (NLTG) and Renewal (NLRG)</b>	€10,220
	<b>Assessment costs</b>	
	Efficacy	€840
	Physicochemical properties of the product	€560
	Criteria with respect to public health	€420
	Criteria with respect to the operator	€840
	Environmental behaviour and environmental fate	€1,540
	Ecotoxicology	€1,960
	Project supervision	€3,080
	Final assessment of the authorisation aspects	€1,400
	<b>Assessment costs for products based on low-risk substances, micro-organisms or pheromones or a comparable substance</b>	
	Efficacy	€1,680
	Physicochemical properties of the product	€140
	Criteria with respect to public health and the operator	€560
	Environmental behaviour and environmental fate	€700
	Ecotoxicology	€700
	Project supervision	€1,960
	Final assessment of the authorisation aspects	€1,400
	<b>Additional fees and charges</b>	
	Each additional active substance	€2,380
	For every 1 to 5 additional uses	€980
	Per tank mix	€2,100
	Additional dRR	€9,520
	Evaluation of summaries and studies concerning physicochemical properties, analytical methods, efficacy, human toxicology and environmental fate and ecotoxicology	Actual costs incurred
	Use of Guidance on honeybees and other pollinators	€3,920
	Use of Guidance on protected cultivation (per 5 crops)	€560
	Use of Guidance on relevance of metabolites in groundwater (B10) - per metabolite	€980
	GAP restriction during application process	€1,120
	Assessment of bridging study, per bridging study	€560
	<b>Additional assessments:</b>	
	Use of Guidance for Birds & Mammals	€1,680
	Use of Guidance for Aquatic organisms	€1,260
	Effect of combination of substances - ecotoxicology	€1,120
	Use of Guidance for non-target organisms	€1,120
	Statistical analysis Complex studies and field studies	€2,240
	Evaluation ecotoxicology model	€7,000
	Human Toxicology assessment import tolerance	€21,000
	Maximum Residue Limit assessment relevant metabolites	€3,640
	<b>Comparative Assessment of crop according to Article 50 Regulation (EC) 1107/2009</b>	
	CA 1: Intake	€840
	Agronomic assessment: per application	€715
	CA 2: Risk assessment	Actual costs incurred

## 1.2 Application to amend current authorisation where NL acts as CMS (NLWTG)

An NLWTG is an application to amend the current authorisation in which the Netherlands is a concerned Member State.

Type of application	Description	Amount
NLWTG	<b>Application to amend current authorisation, the Netherlands is CMS</b>	
	<b>Application costs</b>	€6,720
	<b>Assessment costs</b>	
	Efficacy	€700
	Physicochemical properties of the product	€420
	Criteria with respect to public health	€420
	Criteria with respect to the operator	€560
	Environmental behaviour and environmental fate	€840
	Ecotoxicology	€1,120
	Project supervision	€2,240
	Final assessment of the authorisation aspects	€1,400
	<b>Assessment costs for products based on low-risk substances, micro-organisms or pheromones or a comparable substance</b>	
	Efficacy	€1,680
	Physicochemical properties of the product	€140
	Criteria with respect to public health and the operator	€560
	Environmental behaviour and environmental fate	€700
	Ecotoxicology	€700
	Project supervision	€1,960
	Final assessment of the authorisation aspects	€1,400
	<b>Additional fees and charges</b>	
	Each additional active substance	€2,380
	For every 1 to 5 additional uses	€980
	Per tank mix	€2,100
	Additional dRR	€9,520
	Evaluation of summaries and studies concerning physicochemical properties, analytical methods, efficacy, human toxicology and environmental fate and ecotoxicology	Actual costs incurred
	Use of Guidance on honeybees and other pollinators	€3,920
	Use of Guidance on protected cultivation (per 5 crops)	€560
	Use of Guidance on relevance of metabolites in groundwater (B10) - per metabolite	€980
	GAP restriction during application process	€1,120
	Assessment of bridging study, per bridging study	€560
	<b>Additional assessments:</b>	
	Use of Guidance for Birds & Mammals	€1,680
	Use of Guidance for Aquatic organisms	€1,260
	Effect of combination of substances - ecotoxicology	€1,120
	Use of Guidance for non-target organisms	€1,120
	Statistical analysis Complex studies and field studies	€2,240
	Evaluation ecotoxicology model	€7,000
	Human Toxicology assessment import tolerance	€21,000
	Maximum Residue Limit assessment relevant metabolites	€3,640
	<b>Comparative Assessment of crop according to Article 50 Regulation (EC) 1107/2009</b>	
	CA 1: Intake	€840
	Agronomic assessment: per application	€715
	CA 2: Risk assessment	Actual costs incurred

## 2. Fees and charges mutual recognition plant protection products (WERG)

Type of application	Description	Amount
NLWERGZ, NLWERG	<b>Mutual recognition</b> Mutual recognition can be based on a zonal dossier (NLWERGZ) or on an authorisation under Directive 91/414/EEC (NLWERG) from another Member State	
	<b>Application costs</b>	€4,620
	<b>Assessment costs</b>	
	Efficacy	€700
	Physicochemical properties of the product	€560
	Criteria with respect to public health	Not applicable
	Criteria with respect to the operator	€840
	Environmental behaviour and environmental fate	€2,380
	Ecotoxicology	€2,800
	Project supervision	€2,380
	Final assessment of the authorisation aspects	€1,400
	<b>Assessment costs for products based on micro-organisms, pheromones or comparable active substances, including low-risk substances</b>	
	Efficacy	€1,680
	Physicochemical properties of the product	€700
	Criteria with respect to public health and the operator	€560
	Environmental behaviour and environmental fate	€700
	Ecotoxicology	€700
	Project supervision	€1,960
	Final assessment of the authorisation aspects	€1,400
	<b>Additional fees and charges</b>	
	Each additional active substance	€2,380
	For every 1 to 5 additional uses	€980
	Per tank mix	€2,100
	Additional dRR	€9,520
	Evaluation of summaries and studies concerning physicochemical properties, analytical methods, efficacy, human toxicology and environmental fate and ecotoxicology	Actual costs incurred
	Use of Guidance on honeybees and other pollinators	€3,920
	Use of Guidance on protected cultivation (per 5 crops)	€560
	Use of Guidance on relevance of metabolites in groundwater (B10) - per metabolite	€980
	GAP restriction during application process	€1,120
	Assessment of bridging study, per bridging study	€560
	<b>Additional assessments:</b>	
	Use of Guidance for Birds & Mammals	€1,680
	Use of Guidance for Aquatic organisms	€1,260
	Effect of combination of substances - ecotoxicology	€1,120
	Use of Guidance for non-target organisms	€1,120
	Statistical analysis Complex studies and field studies	€2,240
	Evaluation ecotoxicology model	€7,000
	Human Toxicology assessment import tolerance	€21,000
	Maximum Residue Limit assessment relevant metabolites	€3,640
	<b>Comparative Assessment of crop according to Article 50 Regulation (EC) 1107/2009</b>	
	CA 1: Intake	€840
	Agronomic assessment: per application	€715
	CA 2: Risk assessment	Actual costs incurred

### 3. National amendment application (NLWG)

An NLWG is an application to amend an existing authorisation where only the national addendum needs to be assessed.

Type of application	Description	Amount
NLWG	<b>National application for amendment (national addendum) of an authorisation</b>	
	<b>Application costs</b>	€1,820
	<b>Assessment costs</b>	
	Efficacy	€1,120
	Physicochemical properties of the product	Not applicable
	Criteria with respect to public health	€140
	Criteria with respect to the operator	€140
	Environmental behaviour and environmental fate	€980
	Ecotoxicology	€980
	Project supervision	€1,540
	Final assessment of the authorisation aspects	€1,400
	<b>Assessment costs for products based on micro-organisms, pheromones or comparable active substances, including low-risk substances</b>	
	Efficacy	€1,820
	Physicochemical properties of the product	Not applicable
	Criteria with respect to public health and the operator	€420
	Environmental behaviour and environmental fate	€700
	Ecotoxicology	€700
	Project supervision	€1,960
	Final assessment of the authorisation aspects	€1,400
	<b>Additional fees and charges</b>	
	Each additional active substance	€2,380
	For every 1 to 5 additional uses	€980
	Per tank mix	€2,100
	Additional dRR	€9,520
	Evaluation of summaries and studies concerning physicochemical properties, analytical methods, efficacy, human toxicology and environmental fate and ecotoxicology	Actual costs incurred
	Use of Guidance on honeybees and other pollinators	€3,920
	Use of Guidance on protected cultivation (per 5 crops)	€560
	Use of Guidance on relevance of metabolites in groundwater (B10) - per metabolite	€980
	GAP restriction during application process	€1,120
	Assessment of bridging study, per bridging study	€560
	Additional assessments:	
	Use of Guidance for Birds & Mammals	€1,680
	Use of Guidance for Aquatic organisms	€1,260
	Effect of combination of substances - ecotoxicology	€1,120
	Use of Guidance for non-target organisms	€1,120
	Statistical analysis Complex studies and field studies	€2,240
	Evaluation ecotoxicology model	€7,000
	Human Toxicology assessment import tolerance	€21,000
	Maximum Residue Limit assessment relevant metabolites	€3,640
	<b>Comparative Assessment of crop according to Article 50 Regulation (EC) 1107/2009</b>	
	CA 1: Intake	€840
	Agronomic assessment: per application	€715
	CA 2: Risk assessment	Actual costs incurred

#### 4. Fees and charges, deriving maximal residue limit (MRL)

Type of application	Description	Amount
MRL	<b>Derivation of Maximum Residue Limit (MRL)</b>	
	<b>Application costs</b>	€2,660
	<b>Assessment costs</b>	
	Physicochemical properties of the product	€1,120
	Criteria with respect to public health	€2,240
	Criteria with respect to the operator	€1,960
	Project supervision	€2,660
	Final assessment of the authorisation aspects	€1,400
	<b>Additional fees and charges</b>	
	Evaluation of summaries and studies concerning physicochemical properties, analytical methods and human toxicology	Actual costs incurred

## Chapter 8c – Fees and charges, Plant protection products, other applications

This concerns applications and assessments based on a fixed fee.

1. Fees and charges, other applications for plant protection products where there is no assessment
2. Fees and charges for other plant protection product applications for which additional assessment costs (in addition to the application fee) can be charged if necessary.

### 1. Fees and charges, other applications plant protection products, where there is no assessment

Type of application	Description	Amount
	<b>Administrative changes</b>	
WNT	Change of name authorised product	€280
WNAW	Change of name, address and/or city of the authorisation holder	€280
OT	Transfer of authorised product to a different company, for each company	€280
INTR	Withdrawal of an authorisation for a plant protection product	No fee charged
	<b>Minor changes</b>	
WSGNW	Minor change in the formulation	€910
Add or change product manufacturer, change production location	Change in product production process ( <u>transitional legislation</u> )	No fee charged
NLWATG	Minor change of WG-GA/WG, including restricting scope of permitted use. (for which GAP is not changed and no risk assessment is required)	€280
	<b>Other types of authorisations</b>	
	<b>Derived applications</b>	
AG	New derived authorisation	€910
VAG	Renewal of derived authorisation	€910
UAG	Expansion of derived authorisation	€910
NLWERGV	Administrative renewal of an authorisation obtained by mutual recognition	€910
	<b>Parallel trade applications</b>	
PAG	Parallel trade permit	€910
VPAG	Renewal of parallel trade permit	€910
UPAG	Expansion of parallel trade permit	€910
	<b>Exemption for trial purposes</b>	
PGN	Permit for trial purposes plant protection product, where the treated crops will <b>not</b> be introduced into the food chain	€280
PGR	Permit for trial purposes plant protection product, where the treated crops will be introduced into the food chain, base amount	€350
PGR	Additional amount per 5 crop uses where the treated crop is introduced into the food chain (rules in the table)	€140
VIG	<b>Application for information on animal testing</b>	€560
	<b>Mediation in the exchange of animal test data</b>	Actual costs incurred
TT	<b>Application for approval of an adjuvant</b>	€910
TT-W	<b>Application for changing an adjuvant</b>	€280
EXV	<b>Export declaration</b>	€280
ART38	<b>Exemption as referred to in Article 53 of Regulation (EC) 1107/2009:</b>	Actual costs incurred

**2. Fees and charges for other plant protection product applications for which additional assessment costs (in addition to the application fee) can be charged if necessary**

Type of application	Description	Amount
WYG	Changes to packaging, shelf life or labelling	€420
WYG-AG	Changes to packaging, shelf life or labelling of derived authorisations	€280
PWSG	Change in production process of <u>active substance</u> and/or additional production location of the active substance. Equivalence through Tier I not yet assessed in another Member State	€1,820
PWSGC	Change in production process of <u>active substance</u> and/or additional production location of the active substance. Equivalence has been assessed by another Member State and published on CircaBC.	€280
MTR	Application to derive standard under the Water Framework Directive (derived from AA-EQS/MAC-EQS)	€2,100
IKV	<b>Notification obligation</b> Regulation (EC) No. 1107/2009 Article 56, fourth clause	€280
<b>Assessment Aspect</b>		<b>Amount</b>
	Efficacy	€1,260
	Physicochemical properties of the product	€1,120
	Criteria with respect to public health	€2,240
	Criteria with respect to the operator	€1,960
	Environmental behaviour and environmental fate	€2,520
	Ecotoxicology	€4,900
	Project supervision	€2,660
	Final assessment of the authorisation aspects	€1,400
<b>Additional fees and charges</b>		
	Evaluation of literature (public or proprietary), data (public or proprietary), new information, summaries and studies concerning physicochemical properties, analytical methods, efficacy, human toxicology and environmental fate and ecotoxicology	Actual costs incurred

**Chapter 8d – Annual fee plant protection product**

	Annual fee	Amount
JV	Annual fee plant protection product	€1,635

## Chapter 9a – Fees and charges Biocidal products (BPR)

### General

**Advance payment with subsequent costing** is required for the following types of applications: authorisation of active substances, applications for Union authorisation, applications for simplified authorisation and applications for national authorisation under the BPR, where the Ctgb acts as the evaluating Member State.

With these applications it is difficult to estimate in advance how many hours the assessment will take and there is a large spread in the required assessment time between the applications. The Ctgb invoices for the advance payment at the start of the application process. If necessary, an additional advance payment will be charged.

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

1. Fees and charges for applications for approval of active substances
2. Fees and charges for applications for the authorisation of a product or for the amendment or renewal of a product authorisation, where Ctgb is the evaluating authority (eCA)
3. Fees and charges for applications for the authorisation of a product or for the amendment or renewal of a product authorisation, where the Netherlands is a concerned Member State
4. Fees and charges for other applications or administrative requests

### 1. Fees and charges for applications for approval of active substances (authorisation and renewal of active substances as well as inclusion in Annex I (of Regulation (EU) 528/2012))

Type of application	Application type R4BP	Description	Amount
EU-B	AS-APP, AS-RNL, AS-EVA, AN-APP	Application for authorisation or renewal of authorisation of an active substance, existing or new, or inclusion of an active substance in Annex I of Regulation (EU) 528/2012 This application fee is for a single product type (PT)	€25,000
		When the authorisation of the same substance for multiple PTs is applied for, an additional fee is charged per PT	€5,000
EU-B, TB3EURAP, TB4EURAP	AS-APP, AS-RNL, AS-EVA, AN-APP	Evaluation of a simple dossier or dossier for an application for inclusion of an active substance in Annex I of Regulation (EU) 528/2012	Actual costs incurred with advance payment of €100,000*
EU-B, TB3EURAP, TB4EURAP	AS-APP, AS-RNL, AS-EVA	Average evaluation	Actual costs incurred with advance payment of €200,000*
EU-B, TB3EURAP, TB4EURAP	AS-APP, AS-RNL, AS-EVA	Evaluation of a complex dossier	Actual costs incurred with advance payment of €250,000
<b>Services and activities regarding EU decision-making</b>			
EU-B, TB3EURAP, TB4EURAP	AS-APP, AS-RNL, AS-EVA, AN-APP	Activities regarding European decision-making on applications for authorisation of active substances or inclusion of an active substance in Annex I of Regulation (EU) 528/2012	Actual costs incurred with advance payment of €75,000*

\* Advance payment is based on an average dossier. The advance payment can be changed if justified by the estimated hours. If the advance payment is insufficient to cover all costs, additional costs will be charged.

**2. Fees and charges for applications for the authorisation of a product or for the amendment or renewal of a product authorisation, where the Ctgb is the evaluating authority (eCA).**

Type of application	Application type R4BP	Description	Amount
B-UTN, B-UTH	UA-APP	Application for Union authorisation of biocidal product (new product or re-registration)	Actual costs incurred with advance payment of €70,000*
B-UTFN, B-UFTH	UA-APP	Application for Union Authorisation of biocidal product family (new product or re-registration)	Actual costs incurred with advance payment of €100,000*
B-TN, B-TH	NA-APP	Application for national authorisation of biocidal product (new product or re-registration)	Actual costs incurred with advance payment of €45,000*
B-TFN, B-TFH	NA-APP	Application for national authorisation of biocidal product family (new product or re-registration)	Actual costs incurred with advance payment of €75,000*
B-ET	SA-APP	Application for simplified authorisation of biocidal product	Actual costs incurred with advance payment of €10,000*
B-ETF	SA-APP	Application for simplified authorisation of biocidal product family	Actual costs incurred with advance payment of €17,000*
B-TPR	NA-APP	Application for provisional authorisation of biocidal product	Actual costs incurred with advance payment of €45,000*
B-TPRF	NA-APP	Application for provisional authorisation of biocidal product family	Actual costs incurred with advance payment of €75,000*
B-TR,	NA-RNL	Application for renewal of an authorisation for a biocidal product	Actual costs incurred with advance payment of €45,000*
B-TRF	NA-RNL	Application for renewal of an authorisation for a biocidal product family	Actual costs incurred with advance payment of € 75,000*
B-ETN	SA-APP	Application for renewal of a simplified authorisation for a biocidal product	Actual costs incurred with advance payment of €10,000*
B-ETFN	SA-APP	Application for renewal of a simplified authorisation for a biocidal product family	Actual costs incurred with advance payment of €17,000*
B-GW,	NA-MAC, SA-MAC, UA-MAC	Application for major change in an authorisation for a biocidal product	Actual costs incurred with advance payment of €45,000*
B-GWF	NA-MAC, SA-MAC, UA-MAC	Application for a major change in an authorisation for a biocidal product family	Actual costs incurred with advance payment of €75,000*
B-MRL		MRL biocidal products	Actual costs incurred

\* Advance payment is based on an average dossier. The advance payment can be changed if justified by the estimated hours. If the advance payment is insufficient to cover all costs, additional costs will be charged.

**3. Fees and charges for applications for the authorisation of a product or for the amendment or renewal of a product authorisation, where the Netherlands is Concerned Member State (CMS).**

Type of application	Type R4BP	Description	Amount
B-TWENP, B-TWEHP	NA-MRP	<b>Application for mutual recognition in parallel of a product (not a family)</b>	
		<b>Initial fee</b>	€3,780
		<b>Evaluation fee</b>	€7,560
		<b>Additional fees and charges</b>	
		Comparative assessment	€1,400
		Additional active substance	€700
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SCP	€980
		Additional PT	€980

Type	Type R4BP	Description	Amount
B-TWEFNP	NA-MRP	<b>Application for mutual recognition in parallel of a biocidal product family</b>	
		<b>Initial fee</b>	€5,180
		<b>Evaluation fee</b>	€9,380
		<b>Additional fees and charges</b>	
		Comparative assessment	€1,400
		Additional active substance	€700
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SCP	€980
		Additional PT	€980

Type of application	Type R4BP	Description	Amount
B-TWENS	NA-MRS	<b>Application for sequential mutual recognition of a product (not a family)</b>	
		<b>All-in fee</b>	€11,340
		<b>Additional fees and charges</b>	
		Comparative assessment	€1,400
		Additional active substance	€700
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SCP	€980
		Additional PT	€980

Type of application	Type R4BP	Description	Amount
B-TWEFNS	NA-MRS	<b>Application for sequential mutual recognition of a biocidal product family</b>	
		<b>All-in fee</b>	€14,560
		<b>Additional fees and charges</b>	
		Comparative assessment	€1,400
		Additional active substance	€700
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SCP	€980
		Additional PT	€980

Type of application	Type R4BP	Description	Amount
B-TWER	NA-RNL	<b>Application for renewal of a product authorisation</b>	
		<b>Initial fee</b>	€3,780
		<b>Evaluation fee</b>	€7,560
		<b>Additional fees and charges</b>	
		Comparative assessment	€1,400
		Additional active substance	€700
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SCP	€980
		Additional PT	€980

Type of application	Type R4BP	Description	Amount
B-TWERF	NA-RNL	<b>Application for renewal of the authorisation for a biocidal product family</b>	
		<b>Initial fee</b>	€5,180
		<b>Evaluation fee</b>	€9,380
		<b>Additional fees and charges</b>	
		Comparative assessment	€1,400
		Additional active substance	€700
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SCP	€980
		Additional PT	€980

Type of application	Type R4BP	Description	Amount
B-GWC	NA-MAC	<b>Application for a major change in an authorisation (biocidal product)</b>	
		<b>Initial fee</b>	€3,780
		<b>Evaluation fee</b>	€7,560
		<b>Additional fees and charges</b>	
		Comparative assessment	€1,400
		Additional active substance	€700
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SCP	€980
		Additional PT	€980
B-GWFC	NA-MAC	<b>Major change to an authorisation (family)</b>	
		<b>Initial fee</b>	€5,180
		<b>Evaluation fee</b>	€9,380
		<b>Additional fees and charges</b>	
		Comparative assessment	€1,400
		Additional active substance	€700
		For more than 5 meta-SPCs, an add-on fee is charged for each additional meta-SCP	€980
		Additional PT	€980

#### 4. Fees and charges for other applications or administrative requests for biocidal products for which additional assessment costs (in addition to the application fees) can be charged if necessary

Type of application	Type R4BP	Description	Amount
B-FL	NA-NPF, SA-NPF, UA-NPF	Application to add a product to a biocidal product family	€1,400
B-AW, B-AWF	NA-ADC, NA-TRS, NA-CCL, NA-MRG, SA-ADC, SA-TRS, US-TDC, UA-TRS	Application for administrative change for a single product or biocidal product family in accordance with Implementing Regulation (EU) 354/2013	€280
B-KW, B-KWF	NA-MIC, SA-MIC, UA-MIC	Application for minor change for a single product or biocidal product family * in accordance with Implementing Regulation (EU) 354/2013	€1,260
PB	ET-NOT	Notification of experiment or trial	€560
PAB, VPAB, UPAB,	PP-APP	Application for parallel trade permit	€840
IKV-B	UE-NOT NE-NOT SE-NOT	Application fee for Obligatory Notification biocidal products (reporting new information)	€280
		<b>Assessment costs</b>	
		Evaluation of summaries and studies concerning physicochemical properties, analytical methods, efficacy, human toxicology and environmental fate and ecotoxicology	Actual costs incurred

\* If necessary, assessment costs may be charged

Type of application	Type R4BP	Description	Amount
B-EM	SN-NOT	Notification of simplified authorisation, authorised in another Member State	€280

Type of application	Type R4BP	Description	Amount
B-ST, B-STF	NA-BBS, NABBP, SA-BBP	Application for authorisation of a product or a biocidal product family using the procedure in Implementing Regulation (EU) 414/2013 and (EU) 1802/2016 (Same biocidal product)	
		The standard fee for applications for the same biocidal product is based on an application without a simultaneous administrative change	
		<b>Application costs</b>	€1,400
		<b>Registration product families (per 5 products)</b>	€630
		<b>Additional fees and charges</b>	
		Simultaneous administrative change	€280

## Chapter 9b – Fees and charges Biocidal products (transitional law)

### General

This concerns applications and assessments based on fixed fees. For additional and/or other services and activities for which no fee has been set, the actual costs incurred will be charged.

1. Fees and charges for product applications and other services and activities where one or more existing active substances for the requested PTs are not yet included in the Union list of approved substances or Annex I of Regulation (EU) 528/2012
2. Other or administrative applications

### 1. Fees and charges for product applications and other services and activities where one or more existing active substances for the requested PTs are not yet included in the Union list of approved substances or Annex I of Regulation (EU) 528/2012

Type of application	Description	Amount
TB, UB, WSB, TVB	<b>Application for authorisation, extended authorisation; change/amendment and renewal</b>	
	<b>Application costs</b>	€3,220
	<b>Assessment costs</b>	
	Efficacy	€2,100
	Physicochemical properties of the product	€980
	Criteria with respect to public health and the operator	€1,540
	Environmental behaviour and environmental fate	€2,100
	Project supervision	€2,520
	Final assessment of the authorisation aspects	€1,400
	Assessment costs if comparative assessment is used (fee per aspect)	€1,400
	Evaluation of summaries and studies concerning physicochemical properties, analytical methods, efficacy, human toxicology and environment	Actual costs incurred
KB	<b>Application for establishing frame formulation</b>	€2,800

### 2. Other applications or administrative applications

Type of application	Description	Amount
<b>Other applications</b>	If necessary, assessment costs can be charged for the applications listed below. These costs are estimated in advance.	Actual costs incurred
AB, VAB, UAB	Application for authorisation, renewal or extended authorisation of a derived biocidal product	€840
TVBA	Application for administrative renewal of biocidal product	€840
WWGGA	Application for amendment of WG/GA of an authorised biocidal product (editorial)	€280
WSBNW	Application for a minor change in the formulation of the biocidal product	€840
PAB, VPAB, UPAB	Parallel authorisation of biocidal product, renewal of parallel authorisation of biocidal product, other applications concerning parallel authorisation of a biocidal product	€840
WNT, WNAW, OT	Application for changing the name of an authorised product, changing the name, address and/or city of the authorisation holder, or transferring an authorised product to a different company	€280
WYB	Application to change the packaging or labelling of a biocidal product, other changes (editorial)	€420
EXV	Export declaration	€280
VIB	Request for information on animal testing of biocidal product	€560
	Evaluation of additional efficacy data	Actual costs incurred
	Mediating the exchange of animal testing data	Actual costs incurred
Changing the production location of product under transitional legislation		<b>No fee charged</b>

## Chapter 9c – Annual fee biocidal products

### Annual fee biocidal product

JV	Annual fee	Amount
	<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.</i>	
	Authorisation of a biocidal product	€1,250
	Authorisation of a biocidal product family	€1,060
	Additional charge per family member	€205
	The maximum fee that will be charged for a family is €5,050 per year.	

\*