

Appendix 1. Fees for biocidal active substances and products

A. Fees for evaluation of applications*

* All articles refer to (EU) No 528/2012 (BPR)

Before the Norwegian Environment Agency begins its work with applications or notifications, fees shall be paid. Fees are claimed by the Norwegian Environment Agency by sending out invoices containing the corresponding fee.

If the fee is shown to be higher than the actual costs after finalised assessments (including possible complaint handling), the exceeding fee will be refunded to the applicant for the following application types:

- a) All applications for active substances
- b) Authorisation of biocidal products or biocidal product families with Norway as the reference member state
- c) Union authorisations
- d) Addition if there is a completely new data package for an active substance
- e) Essential use in 180 days (Article 55(1))
- f) Interim authorisation of a product with a new active substance (Article 55(2))

The Norwegian Environment Agency can impose an additional fee for applications mentioned in the second paragraph if the costs clearly exceed the previously imposed fee.

Active substances – fees

Application type	Chemical active substances, fee in NOK	Microorganism as active substances, fee in NOK
Assessment of an active substance	7 333 248	3 666 624
- Addition per extra product-type	3 666 624	1 833 312
Inclusion in Annex I of the BPR (simplified procedure)	3 666 624	1 833 312
Renewal of approval - full evaluation	7 333 248	3 666 624
- Addition per extra product-type	3 666 624	1 833 312
Renewal of approval – limited evaluation	3 666 624	1 833 312
- Addition per extra product-type	1 833 312	915 034

Biocidal products - fees

Application type	Single product, fee in NOK	Product family, fee in NOK
Product authorisation – as reference member state	900 000	1 800 000
- Addition if there is a completely new data package for an active substance	3 666 624	3 666 624
- Addition per extra active substance	72 000	144 000
- Addition for each co-formulant defined as a "Substance of concern"	72 000	144 000
- Addition per extra product-type	72 000	144 000
- Addition per extra user category	72 000	144 000
- Addition for comparative assessment	90 000	180 000
- Addition for evaluation of MRL (Maximum Residue Level)	45 000	90 000
Union authorisation – as reference member state	1 100 000	To be agreed
- Addition per extra active substance	72 000	144 000
- Addition for each co-formulant defined as a "Substance of concern"	72 000	144 000
- Addition per extra product-type	72 000	144 000
- Addition per extra user category	72 000	144 000
- Addition for comparative assessment	90 000	180 000
- Addition for evaluation of MRL (Maximum Residue Level)	45 000	90 000
Mutual recognition	81 120	162 240
- Addition per extra product-type	14 710	29 420
- Addition for each co-formulant defined as a "Substance of concern"	14 710	29 420
- Addition per extra product-type	14 710	29 420
- Addition per extra user category	14 710	29 420
- Addition for comparative assessment	18 333	36 666
Authorisation of same biocidal product	29 420	58 839

Product assessment after simplified procedure		
- Product authorisation – as reference member state (Article 25)	180 000	360 000
- Notification of product (Article 27)	16 224	32 448
Notification of each new product in a product family	16 224	-

Biocidal products – fees for renewals

Application type	Single product, fee in NOK	Product family, fee in NOK
Full assessment – as reference member state	900 000	1 800 000
- Addition per extra active substance	72 000	144 000
- Addition for each co-formulant defined as a "Substance of concern"	72 000	144 000
- Addition per extra product-type	72 000	144 000
- Addition per extra user category	72 000	144 000
- Addition for comparative assessment	90 000	180 000
- Addition for evaluation of MRL (Maximum Residue Level)	45 000	90 000
Full assessment – mutual recognition	81 120	162 240
- Addition per extra product-type	14 710	29 420
- Addition for each co-formulant defined as a "Substance of concern"	14 710	29 420
- Addition per extra product-type	14 710	29 420
- Addition per extra user category	14 710	29 420
- Addition for comparative assessment	18 333	36 666
Limited assessment – as reference member state	300 000	600 000
- Addition for comparative assessment	90 000	180 000
- Addition for evaluation of MRL (Maximum Residue Level)	15 000	90 000
Limited assessment – mutual recognition	45 860	91 720

- Addition for comparative assessment	18 333	36 666
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Biocidal products – fees for changes to authorisations

Application type	Single product, fee in NOK	Product family, fee in NOK
Major change of product authorisation - as reference member state	300 000	600 000
- Mutual recognition	73 441	146 881
Minor change of product authorisation - as reference member state	87 934	175 868
- Mutual recognition	29 203	58 406
Administrative change of product authorisation	14 602	29 203

Biocidal products – other fees

Application type	Single product, fee in NOK	Product family, fee in NOK
Parallel trade permit (Article 53)	58 731	-
Research and development (Article 56)	40 560	-
Essential use in 180 days (Article 55(1))	81 120	-
Interim authorisation of a product with a new active substance (Article 55(2))	146 665	293 330
- Mutual recognition	18 387	36 774

B. Annual fee

The annual fee is a fixed amount for authorised biocidal products and biocidal product families on the Norwegian market.

The annual fee is claimed in accordance with the Norwegian biocides regulation by the Norwegian Environment Agency.

Fixed annual fee	Fee in NOK
Per authorised product	3 120
Per authorised product family	6 240