

In accordance with Article 80 of Regulation (EU) No. 528/2012 of the European Parliament and of the Council and Regulation 25 of S.I. No. 427 of 2013, the following fees shall apply from **1st January 2017**

Fees relating to Biocidal Products

Description	Fee
<u>Notification</u>	
National Notification of a biocidal product	€300
Trivial amendment	€300
Notification under the BPR (Article 17.6 [#] and Article 27 [*])	€300
<u>Biocidal Product Authorisation¹</u>	
Authorisation	€20,000²
Authorisation (mutual recognition)	€2,500
Authorisation (simplified authorisation procedures)	€2,500
Authorisation (the same biocidal product)	€300
Authorisation (Provisional)	€20,000²
Renewal of an Authorisation	€20,000²
Biocidal product family (in addition to the cost of product authorisation)	€20,000²
Additional product-type (in addition to the cost of product authorisation)	€20,000²
<u>Changes to Authorised Products</u>	
Administrative changes	€300
Minor change	€1,500
Major change	€3,000
Minor change (for simplified procedures and mutual recognition)	€1,000
Major change (for simplified procedures and mutual recognition)	€2,000
Union authorisation (major change work)	€50,000
<u>Permits</u>	
Parallel trade permit	€500
Trial permit ('once off' experiment)	€300
Change to a trial permit ('once off' experiment)	€100
Trials permit (experimental/trial programme)	€2,000
Change to a trial permit (experimental/trial programme)	€300
Renewal of trials permit (experimental/trial programme)	€300
Emergency use permit	€1,000
Renewal of an emergency use permit	€300
<u>Annual Registration Fees (ARF) and Other Certificates</u>	
ARF (professional product)	€225*
ARF (non-professional product)	€125*
Re-instating product on the Register/Late ARF (professional product)	€425*
Re-instating a product on the Register/Late ARF (non-professional product)	€225*
Certificate of Inclusion on the Register (Certificate of Free Sale)	€150

* €25 will be transferred to the Poisons Information Centre to cover the cost of maintaining the poisons database

Fees for work on behalf of the European Union (Biocidal Active Substances)

Description	Fee
<u>Active substance evaluation (includes micro-organisms/biologicals)²</u>	
Pre-submission consultations/meetings	€5,000
Dossier receipt, registry and validation check	
Project co-ordination of the evaluation and CAR	€10,000
Examination/peer review of physical and chemical properties and analytical methods	€30,000
Examination/peer review of residues profile	€30,000
Examination/peer review of efficacy profile	€10,000
Examination/peer review of toxicology profile	€65,000
Examination/peer review of environmental fate and behaviour	€85,000
Examination/peer review of ecotoxicology profile	€70,000
Total (single dossier – active substance + one product-type)	€305,000
Additional product-type	€80,000
<u>Article 28 active substance evaluation (Amendment of Annex I)</u>	
Pre-submission consultations/meetings	€2,000
Dossier receipt, registry and validation check	
Project co-ordination of the evaluation and CAR	€6,000
Examination/peer review of physical and chemical properties and analytical methods	€7,000
Examination/peer review of residues profile	€6,000
Examination/peer review of efficacy profile	€7,000
Examination/peer review of toxicology profile	€7,000
Examination/peer review of environmental fate and behaviour	€8,000
Examination/peer review of ecotoxicology profile	€7,000
Total (single dossier – active substance + one representative product)	€50,000
<u>Other Fees relating to substance evaluations</u>	
Co-Rapporteur/eCA fee (for evaluation or peer review)	€50,000
Post-approval submission of confirmatory information/data	€10,000²
Renewal of a substance approval	€185,000³
<u>Product evaluation for Union Authorisation (where Ireland is the evaluating Competent Authority (eCA))¹</u>	
Union Authorisation	€75,000
Biocidal product family (In addition to the cost of Union authorisation)	€20,000
Additional product-type (In addition to the cost of Union authorisation)	€20,000
Renewal of a Union Authorisation	€75,000¹

¹ Additional fees may apply where additional data are required/submitted (e.g. comparative assessment)

² A reduced fee may be applied where a full evaluation is not required.

³ An amended fee may be applied, where appropriate.

(Notification of an additional product that falls under an authorised biocidal product family)

‡ (Notification of a product authorised under simplified procedures in another Member State)