

# **GB BPR active substance application form**

This form is for companies who are:

- making a new application for biocidal active substance approval or renewal in GB
- submitting a biocidal active substance dossier following a successful notification under the GB Review Programme

## Guidance

Please use one form per active substance. Multiple product types can be included in the same application form.

Please [submit completed application forms by email](#).

Please [email HSE's Biocides Delivery Team](#) if you have any questions about filling in this form.

For further guidance on submitting your application visit [HSE Biocides - Active Substances - How to apply](#).

## Formats

Data dossiers should be submitted in IUCLID format.

You should use the relevant GB templates for documents such as the draft Risk Assessment Report (RAR) wherever possible. However, HSE also currently accepts EU versions of such documents.

It is essential that you complete and submit the [UK reference list of studies template](#) for any data you submit. This is to enable HSE to build up an accurate record of the data we hold and must be submitted regardless of any other reference list that exists as part of another template.

You can find out more about what data and supporting information you need to provide in an active substance dossier for the various application types at: [HSE Biocides - Active Substance Approval](#).

## Letters of access

If any part of your application relies on data that are protected under GB BPR, you will need to provide a letter of access from the data owner that gives permission for HSE to use those data.

Letters of access must be valid in the UK. HSE also accepts EU BPR letters of access providing they are accompanied by a written statement from the issuer confirming that the access extends to GB BPR, **and** that the data have been submitted to HSE.

If HSE does not hold the data that your letter of access refers to your application may be delayed or rejected.

## Safety data sheets (SDSs)

SDSs should comply with UK REACH.

## Form sections

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## Section 1 – Application details

### Application type

#### **Section 1 – Application details**

Complete this section for **all** applications.

#### **Application type**

Please mark the relevant box for the type of application you wish to make.

More information about the types of application can be found at: [HSE Biocides - Active Substances - How to apply](#) and [HSE Biocides - Simplified Active Substances](#)

- New active substance approval
- New simplified active substance listing
- Renewal of an active substance approval
- Active substance dossier submission following successful notification under the GB Review Programme

#### **Notification details**

If you are submitting a dossier following successful notification under the GB Review Programme, please provide brief details of the notification. Where relevant you should include the date of the originating open invitation and / or declaration of interest and the date the notification and / or declaration of interest was declared to be compliant.

Click or tap here to enter notification details.

Section 1 – Application details  
Active substance details

**Active substance details**

*Name*

Please provide a name for the active substance according to the International Union of Pure and Applied Chemistry (IUPAC) system. IUPAC names allow for unambiguous identification of substances; however a common name may be provided in addition.

If applicable, this should match the name given on the [GB List of Compliant Notifications](#).

*CAS / EC number*

Please provide the unique numeric identifiers assigned by the Chemical Abstracts System (CAS) and / or European Community (EC) as applicable.

If applicable, these should match the CAS / EC numbers given on the [GB List of Compliant Notifications](#).

<b>Active substance name</b>	<b>CAS number</b>	<b>EC number</b>
Ozone generated from oxygen	-	-

## Section 1 – Application details

### Product types

#### **Product types**

If you are applying for a new active substance approval or renewal or are submitting a dossier following successful notification under the GB Review Programme, please mark the boxes for the product types relevant to your application. Descriptions for each product type are available at: [HSE Biocides - Basics](#).

- PT1 – Human hygiene
- PT2 – Disinfectants and algaecides not intended for direct application to humans or animals
- PT3 – Veterinary hygiene
- PT4 – Food and feed area
- PT5 – Drinking water
- PT6 – Preservatives for products during storage
- PT7 – Film preservatives
- PT8 – Wood preservatives
- PT9 – Fibre, leather, rubber and polymerised materials preservatives
- PT10 – Construction material preservatives
- PT11 – Preservatives for liquid-cooling and processing systems
- PT12 – Slimicides
- PT13 – Working or cutting fluid preservatives
- PT14 – Rodenticides
- PT15 – Avicides
- PT16 – Molluscicides, vermicides and products to control other invertebrates
- PT17 – Piscicides
- PT18 – Insecticides, acaricides and products to control other arthropods
- PT19 – Repellents and attractants
- PT20 – Control of other vertebrates
- PT21 – Antifouling products
- PT22 – Embalming and taxidermist fluids

Section 1 – Application details  
GB Simplified Active Substance List category

***GB Simplified Active Substance List category***

If you are applying for new simplified active substance listing, please indicate which category of the GB Simplified Active Substance List is likely to be applicable for your active substance. More information about the categories can be found at: [HSE Biocides - Simplified Active Substances](#)

- Category A
- Category B

**Now go to:**

- [Section 2 - Company details](#)

## Section 2 – Company details

Applicant

### Section 2 – Company details

Complete this section for **all** applications.

#### *Applicant*

Please provide the applicant details. This is the person, company or taskforce / consortium supporting the active substance. Applicants can be based anywhere in the world.

Where the applicant is a consortium or task force, please provide the consortium / task force name in this section and details of each of the member companies in the section 'Data owners / consortium members'.

If you are part of a consortium or task force and are submitting your data independently of the other members, please provide your details in this section and the name of the consortium / task force in the section 'Data owners / consortium members'.

<b>Company name</b>	The European Ozone Trade Association Limited
<b>Address</b>	EUOTA c/o: Ozone Industries Limited, Unit 3 Regents Court, South Way, Walworth Business Park, Andover, Hampshire, United Kingdom
<b>Postcode</b>	SP10 5NX
<b>Contact name</b>	Mike Prince
<b>Email address</b>	Mike@ozone-industries.co.uk

<b>Company name</b>	EuroO3zon UK Limited
<b>Address</b>	1 St James Court, Whitefriars, Norwich, Norfolk, United Kingdom
<b>Postcode</b>	NR3 1RU

Section 2 – Company details  
Entity acting on behalf of the applicant

<b>Contact name</b>	Mr Bernhard Paolini
<b>Email address</b>	chairman@euro3zon.org

***Entity acting on behalf of the applicant***

If different from the applicant, please provide the details of the person / company that is completing this form and will be the main point of contact for this application. For example, this could be a consultant acting on behalf of a company or the representative of a consortium / task force.

<b>Company type</b>	Consultant
<b>Company name</b>	Exponent International Limited
<b>Address</b>	The Lenz, Hornbeam Business Park, Harrogate, North Yorkshire, UK
<b>Postcode</b>	HG2 8RE
<b>Contact name</b>	Alison McGuire
<b>Email address</b>	amcguire@exponent.com

Section 2 – Company details  
Data owners / consortium members

***Data owners / consortium members***

Where there are data owners in addition to the applicant or the applicant is a consortium / task force, please provide the details of the additional companies in this section.

Where the applicant is part of a consortium or task force and is submitting data independently of the other members, please provide the name of the consortium / task force in this section.

The table can be copied as many times as necessary for the number of data owners / consortium members. You can do this by first clicking or tapping anywhere in the table. A plus sign will appear at the bottom right-hand corner – click or tap on this to add a new data owner / consortium member table.

*Consortium member 10*

<b>Company name</b>	Ozone Industries Ltd.
<b>Address</b>	Unit 3 Regents Court, South Way, Walworth Business Park, Andover, Hampshire, UK
<b>Postcode</b>	SP105NX
<b>Contact name</b>	Mike Prince
<b>Email address</b>	mike@ozone-industries.co.uk

***Invoicing details***

Please provide the details of the person or company that will pay the fees associated with the evaluation of the application.

<b>Company name</b>	The European Ozone Trade Association Limited
<b>Address</b>	EUOTA c/o: Ozone Industries Limited, Unit 3 Regents Court, South Way, Walworth Business Park, Andover, Hampshire, UK
<b>Postcode</b>	SP10 5NX

## Section 2 – Company details

### Invoicing details

<b><i>Contact name</i></b>	Mike Prince
<b><i>Email address</i></b>	Mike@ozone-industries.co.uk

**Now go to:**

- [Section 3 - Checklist](#)

### **Section 3 – Checklist**

You should complete the checklist relevant to the type of application you are making.

By completing the checklist, you are confirming that you understand and accept the requirement to submit the listed documentation.

<b><i>Checklist A – New applications for active substance approval and dossier submissions following successful notification under the GB Review Programme.....</i></b>	<b>13</b>
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### Section 3 – Checklist

Checklist A – New applications for active substance approval and dossier submissions following successful notification under the GB Review Programme

#### ***Checklist A – New applications for active substance approval and dossier submissions following successful notification under the GB Review Programme***

Complete this section if you are making a new application for active substance approval or are submitting a dossier following successful notification under the GB Review Programme.

More specific guidance on the data and documentation required can be found at: [HSE Biocides - Active Substances - How to apply](#).

Please mark the boxes to confirm you understand and accept the requirement to submit the following, as applicable:

- a dossier for the active substance (satisfying the data requirements in Annex II of GB BPR)
- a dossier for the representative biocidal product (satisfying the data requirements in Annex III of GB BPR)
- if the active substance meets at least one of the exclusion criteria listed in Article 5 (1), evidence that Article 5 (2) is applicable
- Safety Data Sheets (SDSs) for the active substance
- Safety Data Sheet (SDS) for the co-formulants of the representative product
- a draft Risk Assessment Report (RAR) for the active substance and representative product
- reference list of studies for the application
- any relevant letter(s) of access
- any pre-submission correspondence with HSE

**Now go to:**

- [Section 4 - Declaration](#)

### Section 3 – Checklist

#### Checklist B – New applications for simplified active substance listing (Category A)

##### ***Checklist B – New applications for simplified active substance listing (Category A)***

Complete this section if you are applying for listing of a simplified active substance in **Category A** of the GB Simplified Active Substance List.

More specific guidance on the data and documentation required can be found at: [HSE Biocides - Simplified active substances](#).

Please mark the boxes to confirm you understand and accept the requirement to submit the following, as applicable:

- evidence to show that the active substance fits within Category A of the GB Simplified Active Substance List
- evidence of a robust consensus of expert opinion or data to show that the substance does not give rise to concern against the criteria in Article 28 (2) of GB BPR
- Safety Data Sheet (SDS) for the active substance, where applicable
- reference list of studies for the application, where applicable
- any pre-submission correspondence with HSE

**Now go to:**

- [Section 4 - Declaration](#)

## Section 3 – Checklist

### Checklist C – New applications for simplified active substance listing (Category B)

#### ***Checklist C – New applications for simplified active substance listing (Category B)***

Complete this section if you are applying for listing of a simplified active substance in **Category B** of the GB Simplified Active Substance List.

More specific guidance on the data and documentation required can be found at: [HSE Biocides - Simplified active substances](#).

Please mark the boxes to confirm you understand and accept the requirement to submit the following, as applicable:

- a dossier for the active substance (satisfying the data requirements in Annex II of GB BPR)
- a dossier for the representative biocidal product (satisfying the data requirements in Annex III of GB BPR)
- Safety Data Sheets (SDSs) for the active substance
- Safety Data Sheet (SDS) for the co-formulants of the representative product
- a draft Risk Assessment Report (RAR) for the active substance and representative product
- reference list of studies for the application
- any relevant letter(s) of access
- any pre-submission correspondence with HSE

**Now go to:**

- [Section 4 - Declaration](#)

## Section 3 – Checklist

### Checklist D – New applications for renewal of an active substance approval

#### ***Checklist D – New applications for renewal of an active substance approval***

Complete this section if you are making a new application for renewal of an active substance approval.

More specific guidance on the data and documentation required can be found at: [HSE Biocides - Active Substances - How to apply](#).

Please mark the boxes to confirm you understand and accept the requirement to submit the following, as applicable:

- the full data dossier that currently supports the approval, including any post-approval data and previous renewals, where relevant
- all relevant data that has been gathered or generated since the initial approval / previous renewal, as applicable
- a draft Risk Assessment Report (RAR) of whether the conclusions of the initial or previous assessment(s) for the active substance remain valid
- reference list of studies for the application

**Now go to:**

- [Section 4 - Declaration](#)

## Section 4 – Declaration

### Section 4 – Declaration

Complete this section for **all** applications.

Physical signing of the declaration is not necessary – providing your name, position / role in the company and the date in this section will validate the declaration.

Application forms without a completed declaration will not be accepted.

#### ***Declaration***

By completing this declaration, I:

- confirm that the information given in this application form is true to the best of my knowledge and belief.
- acknowledge that any incomplete information may delay the processing of my application.
- understand that HSE will send me a link to submit the documentation and data related to my application (as indicated in the relevant checklist and / or by HSE) via the HSE Secure File Sharing Service.

<b>Name</b>	Alison McGuire
<b>Position / role in company</b>	Principal Scientist and Head of Biocides (Europe)
<b>Date</b>	17/11/2025