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**Application form for animal feed trials using FEED ADDITIVES not authorised**

Applicants must complete this form in full, otherwise the application cannot be considered. Please submit the completed application form alongside all supporting documents via email to:

* + For trials in England: [feedadditives@food.gov.uk](mailto:feedadditives@food.gov.uk)
  + For trials in Wales: [regulated.products.wales@food.gov.uk](mailto:regulated.products.wales@food.gov.uk)
  + For trials in Northern Ireland: [nioperationalpolicy@food.gov.uk](mailto:nioperationalpolicy@food.gov.uk)
  + For trials in Scotland: [feedadditivetrials@fss.scot](mailto:feedadditivetrials@fss.scot)

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| **Trial Unique Identifier** |  |

*Unique identifier for the trial (e.g. ‘Name of additive & proposed date of trial’)*

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| **Trial Location** | |
| **Contact name (the person responsible for the trial)** |  |
| **Address** |  |
| **Phone number** |  |
| **Email** |  |

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| **Sponsoring company** | |
| **Contact name** |  |
| **Address** |  |
| **Phone number** |  |
| **Email** |  |

*(Sponsoring company; full contact details for person responsible for sponsoring the trial)*

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| **Local Authority where trial site is located** *(England & Wales only)*  ***To find details on your Local Authority*** [*https://www.gov.uk/find-local-councilhttps://www.gov.uk/find-local-council*](https://www.gov.uk/find-local-council) ***& Trading Standards Office:*** [**https://www.tradingstandards.uk/consumers/support-advice**](https://www.tradingstandards.uk/consumers/support-advice)  ***For trials in Northern Ireland just insert Northern Irelan****d* |  |

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| **Proposed trial start date** |  |

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| **Proposed trial end date** |  |

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| **Name and description of trial additive** |  |

*Including the additive name, the name of the active ingredient and the species and strain of any production organism(s) used) (e.g ‘HEnZime’, a xylanase from E. lefant CBF 600)*

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| **Objective of the trial** |  |

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| **Product from trial in the food chain.** |  |
| Is it proposed that the animals (control or all animals) and/or products from this trial i.e. milk, eggs, enter the food chain? |  |
| Please provide details e.g. control animals only, all animals, products i.e. chicken, eggs, beef, milk. |  |

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| Is it proposed that the animals (control or all animals) at the end of this trial are re-homed? |  |
| Please provide details e.g. number of animals rehomed and all animals or only control animals. |  |

**The following information is required to process your application and must be submitted with the application form:**

|  |  |
| --- | --- |
| Please provide FSA/FSS with evidence that trials are carried out in accordance with Article 3 (2) of Regulation 1831/2003, and the principles and conditions within Regulation 767/2009 or the guidelines set out in Article7(4) of Regulation 1831/2003. If the proposed animals at the end of this trial are intended for the food chain, please provide evidence of no adverse effect on animal health, human health or the environment.  Further a detailed trial (experimental) protocol should include e.g experimental design and procedures,  Contact details- Full contacts details for the sponsoring company, trial site, trial director/site manager, veterinary consultant, feed supplier and analytical laboratory. Water supply, waste contractor.  Animals: Origin, species, category age, gender, breeds, number of animals involved in the trial. Animal management: housing, grouping, separation, stocking density, environmental conditions. Fate of the animals at the end of the trial.  Feeds: Any pre-trial feed, or treatments. Characterisation of the unauthorised feed additive. Relevant authorisations in other countries. Feed ingredients, origin, mixing, composition, form, labelling of test additives and premixtures and feed containing it to prevent them being used for other purposes. Analytical details. Provide certificates of analysis.  Procedure: Control and treatment groups. Numbers of animals per group, selection/replacement/withdrawal parameters. Parameters to be measured, data collection, recording, statistical treatment. Veterinary/welfare inspection, health records, treatments, medication. Procedure for deviations from protocol, and recording unusual incidents. Disposal and recording of unused feed. Compliance with GLP or other standards. Detail if Home Office license required.  Please note that this is not an exhaustive list.  . |  |
| Risk assessment (to cover safety for consumer, target animal, the environment and the worker/user). |  |