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I Acts whose publication is obligatory

- ★ **Commission Directive 2001/79/EC of 17 September 2001 amending Council Directive 87/153/EEC fixing guidelines for the assessment of additives in animal nutrition ⁽¹⁾**

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⁽¹⁾ Text with EEA relevance.

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Acts whose titles are printed in light type are those relating to day-to-day management of agricultural matters, and are generally valid for a limited period.

The titles of all other Acts are printed in bold type and preceded by an asterisk.

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(Acts whose publication is obligatory)

COMMISSION DIRECTIVE 2001/79/EC

of 17 September 2001

amending Council Directive 87/153/EEC fixing guidelines for the assessment of additives in animal nutrition

(Text with EEA relevance)

THE COMMISSION OF THE EUROPEAN COMMUNITIES,

Having regard to the Treaty establishing the European Community,

Having regard to Council Directive 70/524/EEC of 23 November 1970 concerning additives in feedingstuffs⁽¹⁾, as last amended by Directive 2001/46/EC of the European Parliament and of the Council⁽²⁾, and in particular Article 5 thereof,

Whereas:

(1) Council Directive 87/153/EEC of 16 February 1987 fixing guidelines for the assessment of additives in animal nutrition⁽³⁾, as last amended by Directive 95/11/EC⁽⁴⁾, should be amended in the light of advances in scientific and technical knowledge.

(2) It has become apparent that the increasing prevalence of antibiotic-resistant bacteria is of major concern to public health. Resistance caused by the use of antibiotics as feed additives contributes to the overall levels of resistance. The guidelines for additives other than micro-organisms and enzymes should therefore be supplemented by the establishment of a requirement for the dossier to include an assessment of the risk of the selection of and/or transfer of resistance to antibiotics and of any increased persistence and shedding of enteropathogens in order to ensure the safety of the use of those additives. For this purpose, the data required for the risk assessment and methodology to be applied should also be established.

(3) They should be supplemented by the establishment of criteria for the assessment of the risk to the consumer which could result from the consumption of food containing residues of the additive or its metabolites. Based on the residue studies maximum residue limits (MRL's) and withdrawal periods should be established, where appropriate.

(4) The environmental impact of feed additives is important since the additives are normally used over a long period, and the abovementioned guidelines should therefore be supplemented by the establishment of criteria for the assessment of the risk of the additive having an adverse effect on the environment, either directly and or as a result of the effect of products derived from it, whether directly or excreted by the animals into the environment. For the determination of this impact a stepwise approach should be followed based on a first and second phase of studies.

(5) The guidelines should be supplemented by more information on how workers and users can be exposed by the additive. An exposure assessment should be provided in order to take adequate measures.

(6) Confidence in the quality and objectivity of dossiers would be improved if they were supplemented by a critical appraisal of an independent person acknowledged to be an expert in the relevant field. The matters to be assessed in this report should be established in the guidelines.

(7) Experience has demonstrated that the guidelines should be supplemented by more specific criteria concerning the efficacy trials.

⁽¹⁾ OJ L 270, 14.12.1970, p. 1.

⁽²⁾ OJ L 234, 1.9.2001, p. 55.

⁽³⁾ OJ L 64, 7.3.1987, p. 19.

⁽⁴⁾ OJ L 106, 11.5.1995, p. 23.

- (8) Article 9b(1) of Council Directive 70/524/EEC provides that an additive referred to in Article 2(aaa) of that Directive shall initially be authorised for 10 years, after which time, the person holding the authorisation for it may seek renewal of the authorisation for a further period of 10 years. It is necessary to establish guidelines indicating the information which must be included in such an application for renewal and its accompanying dossier.
- (9) Article 9c(3) of Council Directive 70/524/EEC provides that 10 years after a substance has first been authorised, the findings of all or part of the evaluation of data and information contained in the dossier supplied for initial authorisation may be used for the benefit of other persons seeking authorisation to put that substance into circulation. It is therefore necessary to establish guidelines indicating the information which must nonetheless be included in an application and accompanying dossier.
- (10) Scientific and technical knowledge should be taken into account.
- (11) For clarity reasons, it is appropriate to divide the guidelines into those applicable to additives other than micro-organisms and enzymes and those applicable to micro-organisms and enzymes.
- (12) These guidelines have been established on the basis of the *Report of the Scientific Committee on Animal Nutrition on the revision of the guidelines for the assessment of additives in animal nutrition* (adopted on 22 October 1999).
- (13) The measures provided for in this Directive are in accordance with the opinion of the Standing Committee for feedingstuffs,

HAS ADOPTED THIS DIRECTIVE:

Article 1

The Annex to Directive 87/153/EEC is amended as follows:

After its title the text set out in the Annex to this Directive shall be inserted.

Article 2

Member States shall bring into force the laws, regulations and administrative provisions necessary to comply with this Directive by 1 January 2002 at the latest. They shall forthwith inform the Commission thereof.

When Member States adopt those provisions, they shall contain a reference to this Directive or be accompanied by such a reference on the occasion of their official publication. Member States shall determine how such reference is to be made.

Article 3

This Directive shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Communities*.

Article 4

This Directive is addressed to the Member States.

Done at Brussels, 17 September 2001.

For the Commission

David BYRNE

Member of the Commission

ANNEX

PART I

ADDITIVES OTHER THAN MICRO-ORGANISMS AND ENZYMES

GENERAL ASPECTS

This document is intended as a guideline for establishing dossiers on substances and preparations being submitted for authorisation as additives in feedingstuffs or a new usage of an authorised additive. The term 'additive', as used in these guidelines refers to the active chemically specified substances or the preparations containing active substances in the state in which they will be incorporated in premixtures and feedingstuffs. The dossiers must enable an assessment to be made of the additives based on the present state of knowledge and make it possible to ensure their compliance with the fundamental principles laid down for their authorisation, which are the subject of the provisions of Article 3a of Council Directive 70/524/EEC.

Where a dossier concerns an additive consisting of or containing genetically modified organisms within the meaning of Articles 2(1) and (2) of Council Directive 2001/18/EC⁽¹⁾, the dossier must include the additional information specified in Article 7a(1) of Directive 70/524/EEC, in addition to the information required by these guidelines.

The dossiers should include detailed reports of all studies done, presented in the order and with the numbering proposed in these guidelines. They should include references and copies of all published scientific data relevant to the evaluation of the additive. An electronic version of the dossier should be made available. The studies are intended to demonstrate the safety of use of the additive in relation to:

- (a) the target species at the proposed levels of incorporation in the feedingstuff;
- (b) those likely to be exposed to the additive by respiratory, other mucosal, eye or cutaneous contact while handling the additive as such or incorporated into premixtures or feedingstuffs;
- (c) consumers who ingest food products obtained from animals having received the additive, which could contain residues of the additive, or its metabolites; this will generally be ensured by the setting of maximum residue limits (MRLs) and withdrawal periods;
- (d) the animals and the human-beings through the selection and spread of antimicrobial resistance genes;
- (e) the environment arising from the additive itself or by products derived from the additive, either directly and/or excreted by animals.

As a general rule, studies to establish the identity, conditions of use, physico-chemical properties, methods of determination and efficacy of the additive, and also its metabolic fate and residues, physiological and toxicological effects on target species must be provided. When the additive is intended for a category of animals belonging to a defined species, efficacy and residue studies must be performed on this target category. The studies necessary for the evaluation of risks to human health or the environment will depend essentially on the nature of the additive and the circumstances of its use. In this respect, no strict rule is applicable. If necessary, additional information will be requested. Reasons must be given for the omission from the dossier of any data prescribed in these guidelines. In particular, studies of mutagenicity, carcinogenicity and reproduction toxicity studies may only be dispensed with if the chemical composition, practical experience, or other considerations can reasonably exclude these effects.

The studies should be done and reported according to appropriate quality standards (e.g. good laboratory practice (GLP) pursuant to Council Directive 87/18/EEC of 18 December 1986, on the harmonisation of laws, regulations and administrative provisions relating to the application of the principles of good laboratory practice and the verification of their applications for tests on chemical substances⁽²⁾).

⁽¹⁾ OJ L 106, 17.4.2001, p. 1.

⁽²⁾ OJ L 15, 17.1.1987, p. 29.

Expert reports on quality, efficacy and safety should be provided. Their authors, who should have relevant qualifications and be recognised experts in the field concerned, should not have been personally involved in the conduct of the tests included in the dossier. The reports must provide a critical appraisal of the documentation provided by the applicant; a factual summary is not sufficient.

The determination of physico-chemical, toxicological and eco-toxicological properties shall be performed in accordance with the methods established by Council Directive 67/548/EEC of 27 June 1967 on the approximation of laws, regulations and administrative provisions relating to the classification, packaging and labelling of dangerous substances⁽¹⁾, as last amended by Commission Directive 2000/33/EC⁽²⁾, or with updated methods recognised by international scientific bodies. The use of methods other than these must be justified.

Each dossier shall contain an adequate summary, an annex proposal and may contain a monograph. The dossiers relating to antibiotics, coccidiostats and other medicinal substances and to growth promoters must be accompanied by a monograph, conforming to the model provided in Section V, enabling the additive concerned to be identified and characterised in accordance with Article 9n of Directive 70/524/EEC. An identification note conforming to the model in Section VI has to be provided for all additives.

For additives intended exclusively for pet food it may not always be necessary to subject additives to an as exhaustive program of chronic toxicity, mutagenicity, reproductive toxicity and carcinogenicity testing as that required for additives intended for feeding to livestock from which products for human consumption are derived. Residue studies in pet animals are not required.

Study of the metabolic fate of the additive in food producing target animals and in laboratory species used for toxicity testing, is required in order to:

- (a) ensure that there is adequate data on the toxicity of the parent additive and any metabolites produced in the target species to which the consumer might be exposed. To this end a comparison of the metabolic fates of the additive in the target and laboratory animal species used for the toxicity testing is important;
- (b) identify and quantify the appropriate marker residue(s) to be used for setting the MRL for the marker residue and the withdrawal periods for the final product.

⁽¹⁾ OJ L 196, 16.8.1967, p. 1.

⁽²⁾ OJ L 136, 8.6.2000, p. 90.

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1. **Section I: Summary of the data in the dossier**

The summary must follow the order of the guidelines and address all the different parts with reference to the relevant pages of the dossier. It should contain a proposal covering all the conditions for the authorisation sought.

2. **Section II: Identity, characterisation and conditions of use of the additive; methods of control**

2.1. *Identity of the additive*

2.1.1. Proposed proprietary name(s)

2.1.2. Type of additive according to its main function. When possible, evidence of mode(s) of action should be included. Any other uses of the active substance should be specified.

2.1.3. Qualitative and quantitative composition (active substance, other components, impurities, batch-to-batch variation). If the active substance is a mixture of active components, each of which is clearly definable, the main components must be described separately and the proportions in the mixture given.

2.1.4. Physical state, particle size distribution, particle shape, density, bulk density; for liquids: viscosity, surface tension.

2.1.5. Manufacturing process including any specific processing procedures.

2.2. *Characterisation of the active substance(s)*

2.2.1. Generic name, chemical name according to IUPAC nomenclature (International Union of Pure and Applied Chemistry), other generic international names and abbreviations. Chemical abstracts service number (CAS).

2.2.2. Structural formula, molecular formula and molecular weight.

For active substances being fermentation products: microbial origin (name and place of culture collection recognised as an international depository authority, preferably in the European Union, where the strain is deposited, accession number and all relevant morphological, physiological, genetic and molecular characteristics for its identification). For genetically modified strains, information on the genetic modification must be provided.

2.2.3. Purity

Identification and quantification of occurring chemical and microbial impurities and toxic substances, confirmation of the absence of production organisms.

2.2.4. Relevant properties

Physical properties of the chemically specified substances: dissociation constant, pKa, electrostatic properties, melting point, boiling point, density, vapour pressure, solubility in water and organic solvents, K_{ow} and K_{oc} , mass and absorption spectra, NMR data, possible isomers and any other appropriate physical properties.

2.2.5. Manufacturing, purification processes, media used and, for fermentation products, batch-to-batch variation.

2.3. *Characterisation of the additive: Physico-chemical and technological properties*

2.3.1. Stability of each formulation of the additive on exposure to environmental conditions such as light, temperature, pH, moisture, oxygen and packing material. Expected shelf life of the additive as marketed.

- 2.3.2. Stability of each formulation of the additive during the preparation and storage of premixtures and feedingstuffs, in particular stability to anticipated process/storage conditions (heat, moisture, pressure/shear, time and packing material). Possible degradation or decomposition products. Expected shelf life of the additive.
- 2.3.3. Other appropriate physico-chemical or technological properties in order to obtain and keep homogeneous mixtures in premixtures and feedingstuffs, antidusting and electrostatic properties, dispersability in liquids.
- 2.3.4. Incompatibilities or interactions that could be expected with feedingstuffs, carriers, other approved additives or with medicinal products.
- 2.4. *Conditions of use of the additive*
- 2.4.1. Where an additive has significant technological as well as zootechnical effects it has to meet the requirements of both claims. The claims for each additive have to be identified and justified.
- 2.4.2. Proposed technological use in animal feedingstuff manufacture or if appropriate in raw materials.
- 2.4.3. Proposed mode of use in animal nutrition (e.g. animal species or categories and age group/production stage of animal, type of feedingstuff, and contra-indications).
- 2.4.4. Proposed method and level of inclusion in premixtures and feedingstuffs or raw materials if appropriate expressed as proportion of the additive and chemically specified substances by weight for premixtures, for feedingstuffs or raw materials if appropriate, with proposed dose in the final feedingstuff and proposed duration of administration and withdrawal period if appropriate.
- 2.4.5. Data from other known uses of the active substance (e.g. in foodstuffs, human or veterinary medicine, agriculture and industry) must be provided.
- 2.4.6. Proposed material safety data sheet as foreseen by Commission Directive 91/155/EEC⁽¹⁾ defining and laying down the detailed arrangements for the system of specific information relating to dangerous preparations in implementation of Article 10 of Directive 88/379/EEC⁽²⁾ and, if necessary, proposed measures for the prevention of occupational risks and means of protection during manufacture, handling, use and disposal.
- 2.5. *Control methods*
- 2.5.1. Description of the methods used for the determination of the criteria listed under items 2.1.3, 2.1.4, 2.2.3, 2.2.4, 2.3.1, 2.3.2, 2.3.3 and 2.3.4.
- 2.5.2. Description of the qualitative and quantitative analytical methods for routine control of the active substance in premixtures and feedingstuffs. This method has to be validated in a ring test involving at least four laboratories or has to be validated in house following international harmonised guidelines for the in-house validation of methods of analysis⁽³⁾ with respect to the following parameters: applicability, selectivity, calibration, accuracy, precision, range, limit of detection, limit of quantification sensitivity, robustness and practicability. Evidence that these characteristics have been assessed must be made available (2.5.4).
- 2.5.3. Description of the qualitative and quantitative analytical methods for determining the marker residue(s)⁽⁴⁾ of the active substance in target tissues and animal products.

⁽¹⁾ OJ L 76, 22.3.1991, p. 35.

⁽²⁾ OJ L 187, 16.7.1988, p. 14.

⁽³⁾ Method Validation — A Laboratory Guide, Eurachem Secretariat, Laboratory of the Government Chemist, Teddington, United Kingdom, 1996.

⁽⁴⁾ The marker residue is a residue whose concentration is in known relationship with the rate at which the concentration of the total residue in the target tissue depletes to the MRL.

- 2.5.4. The methods mentioned under 2.5.2 and 2.5.3 should be accompanied by information as to the sampling method used, percentage recovery, specificity, accuracy, precision, limits of detection, limits of quantification and validation procedure used. Reference standards of the active substance and/or of the marker residue(s) must be available as well as information on the optimum storage conditions for these reference standards. When devising methods, consideration must be given to the fact that their limits of quantification must be below the MRLs. Moreover, their suitability for routine analysis must be taken into account.

3. **Section III: Studies concerning the efficacy of the additive**

3.1. *Studies on the effects on feedingstuffs*

These studies concern technological additives such as antioxidants, preservatives, binders, emulsifiers, stabilisers, gelling agents, pH modifiers etc., which are intended to improve or stabilise the characteristics of premixtures and feedingstuffs but have no direct biological effect on animal production. All claimed activities or effects of the additive have to be justified by scientific information.

Evidence of the efficacy of the additive must be provided by means of appropriate criteria as reflected in recognised acceptable methods, under the intended conditions of use in comparison with appropriate control feedingstuffs. These investigations must be designed and performed so as to permit a statistical evaluation.

Full information on the active substances, preparations, premixes and feedingstuffs examined, the reference number of the batches, the detailed treatment and testing conditions should be provided. Positive and negative effects, both technological and biological, should be described for each experiment.

3.2. *Studies on the effects on animals*

Studies on zootechnical additives must be performed in target species/animal categories for which the additive is intended in comparison with negative control groups (without antibiotics, growth promoters or other medicinal substances) and, possibly, with groups receiving feedingstuffs containing EU approved additives of known effectiveness used at their recommended dosages (positive control).

Animals used should be healthy and preferably from a homogeneous group.

Studies must permit the evaluation of the efficacy of the additive according to farming practice in the EU. Similar protocol designs should, where possible, be used for all trials so that data can eventually be tested for homogeneity and pooled (if tests so indicate) for statistical evaluation.

No single design is recommended, flexibility being provided to allow for scientific discretion in the design and conduct of the studies. The experimental design used must be justified according to the claim for the use of the additive and must include consideration of adequate statistical power.

3.2.1. For coccidiostats and other medicinal substances

Importance should primarily be attached to evidence of the specific effects (e.g. species controlled, life-cycle stage(s) affected) and particularly to prophylactic properties (e.g. morbidity, mortality, oocyst count and lesion score).

Information on the effect on feed efficiency and liveweight gain should be provided.

The required efficacy data involve three stages of target animal experimentation:

- (a) controlled battery-cage experiments (single and mixed infections);
- (b) controlled floor pen studies (simulated use conditions);
- (c) controlled field trials (actual use conditions).

Simultaneously and when relevant, within trials on efficacy, additional data should be recorded to allow an evaluation of interference with growth and feed conversion (fattening birds, replacement layers and rabbits), effects on egg fertility and hatchability (breeding birds).

3.2.2. For other zootechnical additives

Information should be provided on the effects on feed intake, body weight, feed efficiency (preferably on a dry matter basis), product quality and yield and any other parameter of benefit to the animal, the environment, the producer or the consumer. The studies should include an indication of dose/response relationship where appropriate.

3.2.3. Experimental conditions

Trials should be carried out at least at two different locations. They should be reported individually, giving details of the controls and each experimental treatment. The trial protocol should be carefully drawn up with regard to general descriptive data as follows:

3.2.3.1. Herd or flock: location and size; feeding and rearing conditions, method of feeding; for aquatic species, size and number of tanks or pens at the farm and water quality.

3.2.3.2. Animals: species (for aquatic species intended for human consumption identification shall be made by their colloquial name followed in parenthesis by the latin or Linnean description), breed, age, sex, identification procedure, physiological stage and general health.

3.2.3.3. Number of test and control groups, number of animals in each group. The number of animals involved in the trials must permit statistical analysis. The methods of statistical evaluation used should be stated. At least three independent comparable trials at the level of $p < 0,05$ in each of the claimed animal category(ies) have to be provided to show the effect referred to. In the case of ruminants a lower level of probability could be accepted $p < 0,10$. The report should include all animals or experimental units involved in the trials. Cases which cannot be assessed due to a lack or loss of data should be reported and their distribution within the groups of animals classified.

3.2.3.4. Diets: description of manufacture and quantitative composition of the diet(s) in terms of ingredients used, relevant nutrients (analysed values) and energy. Feed intake records.

3.2.3.5. Concentration of the active substance (and, where that is the case, substances used for comparative purposes) in the feedingstuffs should be established by a control analysis, using the appropriate recognised method. Reference number(s) of the batches.

3.2.3.6. Date and exact duration of testing. Date and nature of the examinations performed.

3.2.3.7. Dose determination studies: the purpose of these studies is to explain the rationale for the selection of a dose or dose range claimed to be optimally effective. Dose determination will be based on a control (without antibiotics, growth promoter or other medicinal substances) and at least three non-zero levels in target animals.

3.2.3.8. The timing and prevalence of any undesirable consequences of treatment in individuals or groups must be reported (give details of the observation programme used in the study).

3.2.3.9. All additives studied under farm conditions must have good scientific evidence of safety for the user, consumer, animal and the environment. Where an additive does not meet the requirements for consumer safety any study undertaken should be designed to prevent animal products derived from the test animals from entering the human food chain.

3.3. *Studies on the quality of animal produce*

Animal products should be examined for organoleptic, nutritional, hygienic and technological qualities as appropriate.

3.4. *Studies on the effects on the characteristics of animal wastes*

If the additive is intended to modify some characteristics of animal waste (e.g. nitrogen, phosphorus, odour, volume), then studies demonstrating these properties are required.

4. **Section IV: Studies concerning the safety of use of the additive**

The studies outlined in this section are intended to permit assessment of:

- the safety of use of the additive in the target species,
- any risk associated with the selection and/or transfer of resistance to antibiotics and increased persistence and shedding of enteropathogens,
- the risks to the consumer which could result from the consumption of food containing residues of the additive or its metabolites,
- the risks from respiratory, other mucosal, eye or cutaneous contact for persons likely to handle the additive as such or as incorporated into premixtures or feedingstuffs,
- the risks of adverse effects on the environment, from the additive itself or by products derived from the additive, either directly and/or excreted by animals.

Consideration should be given to known incompatibilities and/or interactions between the additive and veterinary medicines and/or components of the diet relevant to the species concerned.

These studies will normally be required in their entirety for each additive unless a specific exclusion or modification is specified in the Directive.

A more limited submission will normally be accepted for a proposed extension of the authorised use to a species which is physiologically and metabolically close to one in which the use of the additive has already been approved. This reduced data set should demonstrate safety to the new species and lack of significant differences in metabolic fate and residues in edible tissues. The proposed MRL and withdrawal period for the species must be justified.

In order to assess the risks for the consumer and consequently the determination of the MRLs and the withdrawal period the following information has to be provided:

- the chemical structure of the active substance,
- the metabolism in the proposed target species,
- the nature of the residues in these target species,
- tissue depletion study of residues,
- data on the biological effects of the active substance together with its metabolites.

A knowledge of the bioavailability of the residues (both unbound and bound) may also be useful namely when many metabolites are produced and no marker residues is evidenced (see section 4.1.3.3).

Furthermore, knowledge of the composition and of the physico-chemical and biological properties of the major excreted materials deriving from the additive are required to define the extent of the studies necessary for assessment of the risk of adverse effects on the environment or persistence in the environment (see paragraph 4.5).

4.1. *Studies on target species*

4.1.1. Tolerance tests on target species/animal categories

The aim is to determine a safety margin (i.e. margin between the maximum proposed dose-level in feedingstuffs and the minimum level resulting in unfavourable effects). However a safety margin of a factor of at least 10 is considered sufficient to require no further testing. Such a tolerance test must be conducted in the target species/animal categories preferably through the entire length of the production period although a test period of one month would normally be acceptable. This requires at least the assessment of clinical signs and other parameters to ascertain effects on target animal health. A negative control group (without antibiotics, growth promoter or other medicinal substances) has to be included. Depending on the toxicological profile additional parameters may also be required. Any adverse effects detected during efficacy trials should also be reported in this section.

Whenever the product is intended for use in animals which may be used for breeding, studies should be conducted to identify possible impairment of male or female general reproductive function or harmful effects on progeny resulting from the administration of the additive under investigation.

4.1.2. Microbiological safety of the additive.

4.1.2.1. All studies should be performed with the highest proposed dose level

4.1.2.2. If the active substance possesses antimicrobial activity at feed concentration level, the minimum inhibitory concentration (MIC) should be determined in appropriate pathogenic and non-pathogenic, endogenous and exogenous bacteria, according to standardised procedures.

4.1.2.3. Tests to determine the ability of the additive to:

- induce cross-resistance to relevant antibiotics,
- select resistant bacterial strains under field conditions in the target species and, if so, investigations on the genetic mechanisms for transfer of the resistance genes.

4.1.2.4. Tests to determine the effect of the additive:

- on a number of opportunistic pathogens present in the digestive tract (e.g. *enterobacteriaceae*, *enterococci* and *clostridia*),
- on the shedding or excretion of relevant zoonotic micro-organisms, e.g. *salmonella* spp, *campylobacter* spp.

4.1.2.5. In cases where the active substance shows an antimicrobial action, field studies to monitor for bacterial resistance to the additive should be provided.

4.1.3. Metabolism and residue studies

4.1.3.1. The aim of the studies is to:

- establish the metabolic pathways of the active substance as a base for its toxicological evaluation,
- identify residues and establish their kinetics in the edible tissues and products (milk, eggs),
- identify the excreted substances as a prerequisite for assessing their impact on the environment.

Occasionally, e.g. for fermentation-derived additives, it could be necessary to extend these studies to other substances added to or derived during the fermentation process. Possession of a toxicity significant in relation to that of the active component(s) of the additive would exemplify this circumstance.

4.1.3.2. Pharmacokinetics

The planning and experimental design of the studies must take into account the anatomical, physiological (age, type, sex), zootechnical category and environmental peculiarities of the target population. When appropriate, the influence of the intestinal or ruminal microflora, enterohepatic circulation or caecotrophy must be considered. The dose regimen tested must be that intended for use, and possibly a multiple of that dose if justified. The active substance (including the labelled substance) must be incorporated into the feed unless there is justification for not doing so.

The studies required are the following:

- metabolic balance and kinetics in the plasma/blood following a single dose administration in order to assess the rate and extent of the absorption, distribution and excretion (urine, faeces, gills, bile, expired air, milk or eggs),
- identification of the major (> 10 %) metabolites in the excreta; except if a minor (< 10 %) metabolite would appear to be of toxicological concern,
- distribution of labelled material in tissues and products following a single dose administration to animals already in steady state equilibrium achieved with unlabelled additive.

The studies mentioned in 4.1.3.1 and 4.1.3.2 should include isotope tracer or alternative relevant methods.

4.1.3.3. Study of the residues

- identification of those residues (parent compound, metabolites, degradation products, bound residues⁽¹⁾) which represent more than 10 % of the total residue (except if a minor metabolite would appear to be of toxicological concern) in the edible tissues and products (milk, eggs) at metabolic equilibrium, i.e. following multi-dose administration of the labelled substance; ratio of the marker residue to total residues,
- kinetic study of the residues in the tissues (including milk and eggs when appropriate) during the depletion period following achievement of steady-state and using the highest level proposed metabolic profiling, identification of the target tissue⁽²⁾ and of the marker residue,
- depletion study of the marker residue from the target tissues (including milk and eggs if appropriate) after withdrawal of the additive following its repeated administration according to the proposed conditions of use and sufficient to have reached steady-state, in order to set a withdrawal period on the basis of the fixed MRL,
- the withdrawal period for the additive must not be less than the time necessary for the concentration of the marker residue determined in the target tissue to fall below the MRL value (95 % confidence limit). Spaced time points, suitably chosen by reference to the depletion phase of the active substance and its metabolites, and at least four animals per point depending on the species (size, genetic variability) should be considered as a minimum requirement⁽³⁾.

⁽¹⁾ Bound residues correspond to the tissue residual fraction that is not extractable using physico-chemical or biological means. They derive from the covalent binding of a metabolite of the compound with cellular macromolecules.

⁽²⁾ The target tissue is the edible tissue selected to monitor for the total residue in the target animal.

⁽³⁾ For the determination of a withdrawal period, the suggested minimum numbers of healthy animals sampled at each slaughter or time point are the following:

- lactating cattle, eight, including animals at second or subsequent lactations (four high yielding cattle at an early stage of lactation and four low yielding cattle at a late stage of lactation),
- other large animals, four per sampling time,
- poultry, six per sampling time,
- laying birds, ten eggs per time point,
- fish, ten per sampling time.

4.2. *Studies on laboratory animals*

These studies must be carried out with the active substance using internationally recognised standard test methods as described in the OECD Guidelines for methodological details or in Directive 67/548/EEC, and according to the principles of good laboratory practice (GLP). Additional studies on particular metabolites produced by the target species may be necessary if these are not formed to a significant extent in the laboratory test species. Also where there is data in man this may need to be taken into consideration in deciding what additional studies to conduct.

4.2.1. Acute toxicity

Acute oral toxicity studies should be carried in at least two mammalian species. One laboratory species may be replaced, if appropriate, by a target species. It will not be necessary to identify a precise LD₅₀; an approximate determination of the minimum lethal dose is normally adequate. In order to reduce the number and suffering of the animals involved, the maximum dosage should not exceed 2 000 mg/kg body weight and alternative methods (limit test, fixed dose method, acute toxicity class method) are recommended.

Risks to workers should be assessed in a series of studies using the product (active substance plus carrier in the form in which it is to be made available commercially). Studies on skin irritancy must be performed and if these give positive results, mucous membrane (e.g. eye) irritancy should be assessed. Allergic potential — skin sensitisation potential should also be assessed. Acute inhalation studies should be performed if the product is likely to form a respirable dust or mist.

4.2.2. Genotoxicity studies including mutagenicity

In order to identify active substances and, if appropriate, their metabolites and degradation products with mutagenic and genotoxic properties, a selected combination of at least three different genotoxicity tests must be carried out. The test battery should normally include prokaryotic and eukaryotic systems tests including mammalian *in vitro* and *in vivo* tests systems. If appropriate the tests should be performed without and with mammalian metabolic activation.

Reasons for the choice of the tests with regard to their reliability to assess genotoxic effects on different genetic endpoints at the gene, chromosome and genome level should be given. Additional tests may be indicated depending on the outcome of the tests and taking into consideration the whole toxicity profile of the substance as well as the intended use. Tests must be carried out according to established and up-to-date validated procedures. When the test target is bone marrow, proof of exposure of the cells to the test substance is required in the case of a negative result.

4.2.3. Subchronic (90-day) oral toxicity studies

The duration of the tests must be at least 90 days. For additives intended for use in food producing animal species the studies should be carried out on two animal species, of which one should be a non-rodent species, which may be the target species. For additives intended for use in animals not for human consumption the studies on the target species are sufficient: the active substance must be administered orally at least at three levels in addition to a control group to obtain a dose response.

The maximum dose should normally reveal evidence of harmful effects. The lowest dose level should not produce any evidence of toxicity.

4.2.4. Chronic oral toxicity studies (including carcinogenicity studies)

A chronic toxicity study, which may include examination of carcinogenicity, must be carried out in at least one rodent species.

Carcinogenicity studies may not be necessary if the active substance and its metabolites:

- give consistently negative results in an appropriate range of genotoxicity tests,
- are not structurally related to known carcinogens, and
- give no effects indicative of potential (pre)neoplasia in chronic toxicity assays.

4.2.5. Reproduction toxicity studies including teratogenicity

4.2.5.1. Two generation reproduction toxicity study

- Studies of reproductive function must be carried out and extend over at least two filial generations (F1, F2) and may be combined with a teratogenicity study. The substance under investigation shall be administered to males and females at an appropriate time prior to mating. Administration should continue until the weaning of the F2 generation.
- All relevant fertility, gestation, parturition, maternal behaviour, suckling, growth and development of the F1 offspring from fertilisation to maturity and the development of the F2 offspring to weaning must be carefully observed and reported.

4.2.5.2. Teratogenicity study

The teratogenicity study covers embryo and foetotoxicity. It must be carried out in at least two species.

4.2.6. Metabolism and disposition studies

Studies on absorption, distribution in the body fluids and tissues, excretion routes, must be performed. A metabolic study including the metabolic balance and identification of the main metabolites in the urine and faeces should be performed in animals of both sexes and the same strains as those used in the toxicological studies. A single dose of the labelled molecule (see 4.1.3) should be administered at steady state equilibrium reached using the unlabelled compound at a dose similar to the highest level proposed for use in the target animal.

4.2.7. Bioavailability of residues

The assessment of the risk to consumers related to certain residues contained in animal products, namely bound residues, may take into account an additional safety factor based on the determination of their bioavailability using appropriate laboratory animals and recognised methods.

4.2.8. Other specific toxicological and pharmacological studies

Further studies providing additional information useful for the assessment of the safety of the active substance and its residues should be conducted if there is any reason for concern.

4.2.9. Determination of a no observed effect level (NOEL)

All the above findings together with all the relevant published data (including any suitable information on effects of the active substance in human) and information, where appropriate, on closely related chemical structures should be taken into consideration in identifying a NOEL expressed as mg/kg body weight per day. The lowest NOEL should be selected.

However the NOEL to be used for the calculation of the ADI should be selected on the basis of toxicological or pharmacological effects as appropriate. For some additives, e.g. antibacterials, an ADI may be better established on the basis of effects on the human gut microflora. In the absence of internationally accepted and validated methods for describing gut flora, effects on selected and sensitive human gut bacterial strains may be more appropriate.

4.3. *Safety evaluation for the human consumer*

4.3.1. Proposal of the acceptable daily intake (ADI) for the additive

An ADI should where appropriate be proposed.

The ADI (expressed as mg of additive or additive related material per person per day) is derived by dividing the NOEL mg/kg body weight by an appropriate safety factor and multiplying by a mean human body weight of 60 kg. This NOEL expressed as mg/kg body weight per day may be selected using toxicological or pharmacological findings. In some cases an ADI based on the additives microbiological properties may be more relevant. The choice will depend on which property is most relevant in terms of health hazard to the consumer.

The safety factor used to determine the ADI for a particular additive should be selected, bearing in mind the following:

- the nature of the biological effect used to identify the NOEL,
- the relevance of this effect to man and the reversibility of the effect,
- the range and quality of the data used to identify the NOEL,
- any knowledge of the effect(s) of the residue constituents.

It is customary to employ a safety factor of at least 100 in calculating the ADI (i.e. a factor of ten to allow for potential interspecies variation and a further factor of ten to allow for possible differences in response between individual humans). When data on the active substance are available for human beings a lower safety factor may be acceptable.

4.3.2. Proposal of the maximum residue limits (MRLs) of the additive

It is assumed in calculating the MRL that the intake of edible tissue, milk and egg products is the sole source of potential human exposure. If this is not the case an allowance must be made for other sources.

A number of these substances have been used as feed additives and for other applications. In such cases the calculated MRLs would be expected to be the same. There may also be instances where on strict scientific considerations different MRLs are calculated for each use when the route, amount, dosage frequency and duration of dosing, are sufficiently different from those appropriate to use as feed additive, that there are evidences indicating that the kinetics and/or metabolism may result in different residue profiles. In such circumstances it is anticipated that the strictest MRL will be applied.

To establish an MRL the chemical nature of the drug-related material which is intended to be used to specify the tissue residue levels must be defined. This is termed the marker residue. This residue constituent must not necessarily be the toxicologically relevant residue but has to be chosen as a suitable indicator to represent the total significant residue. The ratios of the marker residue/total residues in connection with the ADI (i.e. ratio of the marker residue/total radioactive residues, marker residue/all biologically active residues) should be established at all the time points during the depletion studies. In particular, this ratio should be known at the time point retained to elaborate MRLs. A suitable analytical method for this marker residue must also be available to ensure compliance with the MRL.

In establishing MRLs (expressed as g/kg of marker residue per kg of edible wet tissue or product) on the basis of an ADI, the following daily human food consumption values should be applied:

	Mammals	Birds	Fish
Muscle	300 g	300 g	300 g (*)
Liver	100 g	100 g	
Kidney	50 g	10 g	
Fat	50 g (**)	90 g (***)	
+ Milk	1 500 g		
+ Egg		100 g	

(*) Muscle and skin in natural proportions.

(**) For pigs 50 g of fat and skin in natural proportions.

(***) Fat and skin in natural proportions.

The individual MRLs in different tissues should reflect the depletion kinetics of the residues within those tissues in the animal species intended for use. An analytical method with a limit of quantification below the MRL is required (see section II point 2.5.3).

If a substance could result in a residue in tissues and produce, the MRLs should be proposed in such a way that the total amount of toxicologically (or microbiologically) significant residue ingested⁽¹⁾ daily should be lower than the ADI (see table above).

The MRL should be set only after consideration and inclusion of any other potential sources of exposures of the consumer to components of the residues.

For certain additives residues could arise below the MRL values in milk, eggs or meat which could nonetheless interfere with food quality in particular food processing procedures, e.g. use of milk in cheese making. For such additives, it may be appropriate to consider a 'maximum (food product) processing compatible residue' in addition to establishing MRL values.

There are some circumstances where a MRL will not be required such as:

- no bioavailability of the residues and no harmful effect on the human gut including its microflora,
- complete degradation to nutrients or harmless substances in the target species,
- ADI 'not specified' because of low toxicity in animal tests,
- where use is restricted entirely to feed for pet animals,
- where a substance is also approved as a food additive⁽²⁾, a MRL is normally not required if the marker residue is primarily the parent substance and it constitutes only an insignificant fraction of the ADI of the food additive.

4.3.3. Proposal of the withdrawal period for the additive

The withdrawal period will be set on the basis of the MRLs. The withdrawal time comprises the period after cessation of the administration of the proposed formulation of the additive which is necessary to enable the residue levels to fall below the MRLs (95 % confidence limit).

⁽¹⁾ Proposed calculation: (500 g meat (consisting of 300 g muscle, 100 g liver, 50 g kidney, 50 g fat) or 500 g poultry (consisting of 300 g muscle, 100 g liver, 10 g kidney, 90 g fat) or 300 g fish) + 1 500 g milk + 100 g egg.

⁽²⁾ In accordance with Council Directive 89/107/EEC of 21 December 1988 on the approximation of the laws of the Member States concerning food additives for use as foodstuffs intended for human consumption (OJ L 40, 11.2.1989, p. 27).

To establish a withdrawal period a particular edible tissue may be identified as a surrogate for others, often termed the target tissue.

4.4. *Worker safety assessment*

Workers can be exposed mainly by inhalation or topical exposure while manufacturing or handling or using the additive e.g. farm workers are potentially exposed when handling or mixing the additive. Additional information on how the substances are handled should be provided. An assessment of risk to workers should be included.

Experience in the manufacturing plant is often an important source of information in evaluating the risks to workers from exposure to the additive itself by both airborne and topical routes. Of particular concern are additives/additive-treated feed and/or animal excreta, which are in or may give rise to, a dry powdery form and feed additives which may have allergenic potential.

4.4.1. Toxicological risk assessment for worker safety

4.4.1.1. Effects on the respiratory system

Evidence should be provided that airborne levels of dust will not constitute a hazard to the health of workers. This evidence should include where necessary: inhalation tests in laboratory animals, published epidemiological data and/or the applicants own data on its workplant and/or irritancy and respiratory system sensitisation tests.

4.4.1.2. Effects on the eyes and skin

Where available, direct evidence of absence of irritancy and/or sensitisation should be provided from known human situations. This should be supplemented by findings from validated animal tests for skin and eye irritation, and for sensitisation potential using the appropriate additive.

4.4.1.3. Systemic toxicity

The toxicity data generated to meet safety requirements (including repeated dose toxicity, mutagenicity, carcinogenicity and reproductive testing) should be used to assess other aspects of worker safety. In doing so, it needs to be remembered that contamination of skin and/or inhalation of the additive are the most likely routes of exposure.

4.4.2. Exposure assessment

Information should be provided on how the use of the additive is likely to give rise to exposure by all routes — inhalation, through the skin or by ingestion. This information should include a quantitative assessment, where available, such as typical airborne concentration, dermal contamination or ingestion. Where quantitative information is not available, sufficient information should be given to enable an adequate assessment of exposure to be made.

4.4.3. Measures to control exposure

Using the information from the toxicology and exposure assessment, a conclusion should be drawn about the risks to health of the users (systemic, toxicity, irritancy or sensitisation) when using measures to control exposure, which are reasonable, in the circumstances. If the risk is unacceptable, then precautionary measures should be taken to control or to eliminate exposure. Product reformulation or modification of the procedures for production, use and/or disposal of the additive are preferred solutions. Use of personal protective devices should only be regarded as a measure of last resort to protect against any residual risk once control measures are in place.

4.5. *Environmental risk assessment*

Consideration of the environmental impact of feed additives is important since the administration of feed additives is typically over a long period (even for lifetime), large groups of animals may be involved and many additives are poorly absorbed and therefore excreted intact to a considerable extent. Nonetheless, in a number of cases, the need for environmental assessment may be limited. It is inappropriate to set strict rules in this general guideline. To assist in determining the environmental impact of a feed additive, a stepwise approach should be followed (see decision tree), where in the first phase, additives which do not need further testing can be clearly identified. For other additives a second phase of studies (Phase IIA) are needed to provide additional information, based upon which further studies (Phase IIB) may be considered necessary. Studies, when applicable, should be conducted according to Council Directive 67/548/EEC.

4.5.1. Phase I assessment

The purpose of Phase I assessment is to determine whether or not a significant environmental effect of an additive or its metabolites is likely, based largely on data already established for other purposes.

Exemption from Phase II assessment may be made on one of two criteria:

- (a) The chemical nature and the biological effect of the additive and its use indicate that impact will be negligible: i.e. where the additive and/or its main (more than 20 % of the total residues in the excreta) metabolite(s) are:
 - physiological/natural substances (e.g. a vitamin, or a mineral) that will not alter the concentration in the environment, unless there is evident reason for concern (e.g. copper),
 - additives intended for companion animals (excluding horses).
- (b) The worst case predicted environmental concentration (PEC) is too low to be of concern.

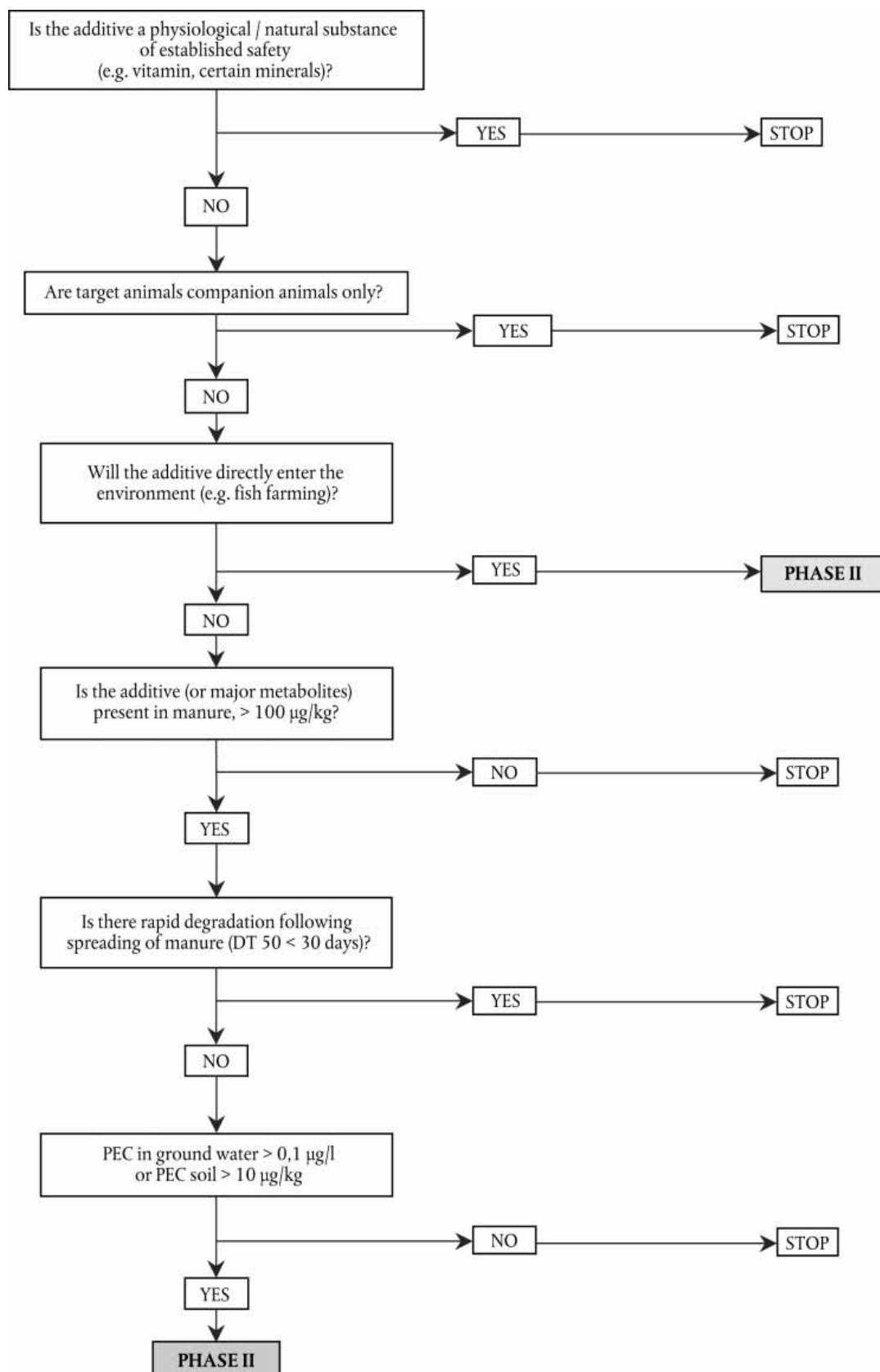
The worst case PEC for soil is likely to arise as a result of manure generated during maximum excretion of the major residue constituents (the additive and/or its major metabolites), being spread on land. The PEC should be evaluated for each major residue constituent in manure and for each compartment of concern. For the terrestrial compartment, if the PEC does not exceed 100 µg/kg for the sum of the major residue constituents in manure, or if the major residue constituents in manure are readily degraded (degradation time DT 50 < 30 days) (in case these data are available) to natural constituents or to concentrations of less than 100 µg/kg, or if the PEC in soil (5 cm depth) is less than µg/kg, then no further assessment is required.

The worst case PEC for water may arise either from direct transfer of spilled feed or excreta containing the additive and its metabolites into water bodies or from the leaching of material in excreta or soil into ground water. If the PEC for contamination of water bodies or ground water is reliably estimated to be less than 0,1 µg per litre no Phase IIA assessment of the environmental impact of the additive on the aqueous compartment is necessary.

If the applicant cannot demonstrate that the proposed additive falls into any of these exemption categories or when the additive is directly released in the environment (e.g. aquaculture), Phase II assessment will normally be required.

ENVIRONMENTAL RISK FROM FEED ADDITIVES

Decision tree Phase I



4.5.2. Phase II assessment

Phase II assessment is in two parts: Phase IIA and Phase IIB.

The bioaccumulation potential of the additive and/or its main metabolites and its influence on the predicted safety margin should be assessed. Bioaccumulation is not considered to be potentially significant if e.g. K_{ow} (partition coefficient) is < 3 . Appropriate Phase IIB studies will generally be needed if such safety margins cannot be established.

4.5.2.1. Phase IIA

The purpose of Phase IIA assessment is to identify risk to the environment by:

- refining the calculation of the PEC(s),
- determining the relationship between exposure, the levels of additive and/or major metabolites and short term adverse effects in relevant surrogate animal and plant species for the environmental compartment(s) of concern,
- using these findings to determine the predicted no effect concentration(s) (PNEC) value(s).

The following sequential procedure is recommended to determine the risk:

- (a) If not already completed in Phase I, a more refined PEC should be calculated for each environmental compartment of concern. In ascertaining the PEC account should be taken of:

- the concentration of additive and/or its major metabolites in manure following administration of the additive to animals at the proposed dose level. This calculation should include consideration of excreta volumes and dosage rates,
- the potential dilution of the excreted additive related material due to normal manure processing practice and storage prior to its application to land,
- the adsorption/desorption of the additive and its metabolites onto soil, the persistence of residues in soil (DT_{50} and DT_{90}); sediment in case of aquaculture,
- other factors such as photolysis, hydrolysis, evaporation, degradation in soil or water sediment systems, dilution through ploughing etc.

The highest value for the PEC obtained by these calculations for each environmental compartment of concern should be adopted for level IIA risk assessment purposes.

If a high persistence in soil ($DT_{90} > 1$ year) at concentrations in excess of 10 g/kg soil is anticipated at steady state, a level IIB assessment may be needed.

- (b) The levels producing serious short-term adverse effects for various trophic levels in the environmental compartments of concern (soil, water) must next be determined. These tests should follow OECD⁽¹⁾ or similar well-established guidelines. Suitable tests for the terrestrial environment include: toxicity to earthworms (50 % lethal concentration, LC_{50} value), phytotoxicity (50 % effective concentration, EC_{50} value) in terrestrial plants, effects on soil micro-organisms (e.g. EC_{50} for effects on methanogenesis and nitrogen fixation). For the aquatic environment: fish: a 96-hour LC_{50} study; *Daphnia magna*: a 48-hour EC_{50} study; algae: an LC_{50} study and a toxicity study for sediment organisms.
- (c) Calculation of the PNEC value for each compartment of concern should be carried out. This is derived normally by taking the lowest value observed (i.e.: the result in the most sensitive species) for an adverse effect in the above ecotoxicity tests and dividing by a safety factor of at least 100 depending on the indicator and number of test species used.
- (d) The calculated PEC and PNEC values should be compared. The acceptable ratio of the PEC to the PNEC value will depend on the nature of the test result used to determine the PNEC. Normally it will be between 1 and 0.1. If significantly lower ratios than these are identified, further ecotoxicological tests are unlikely to be necessary unless bioaccumulation is expected. Conversely, higher ratios will require some Phase IIB testing.

⁽¹⁾ OECD Guidelines for Testing of Chemicals.

4.5.2.2. Phase IIB (more detailed toxicological studies)

For those additives where, following Phase IIA assessment, doubt remains regarding their environmental impact, more detailed studies are required of the effects on biological species in the environmental compartment(s) in which Phase IIA studies indicate possible concern. In this situation, further tests are needed to determine the chronic and more specific effects on appropriate animal, plant and microbial species. It may be that in the Phase IIA assessment the PEC value has been over estimated. To demonstrate this it may be necessary to carry out measurements of the environmental concentrations and, of the persistence of the additive and/or its major metabolites in field use situations.

Suitable additional ecotoxicity tests are described in a number of publications, e.g. in OECD guidelines. Three categories of environmental species may need to be considered, animals, plants and micro-organisms. Careful choice of such tests is necessary to ensure that they are appropriate to the situation in which the additive and/or its metabolites may be released and dispersed in the environment.

The assessment of the impact on the terrestrial compartment may include:

- a sublethal study of the effects on earthworms, further studies of the impact on soil microflora, phytotoxicity tests on a range of economically important plant species, studies on grassland invertebrates including insects and feral birds,
- NB: a separate evaluation of mammalian toxicity may not be necessary, as this aspect is likely to be addressed by mammalian toxicity testing conducted to determine the ADI.

The assessment of the impact on the aquatic compartment may:

- include chronic toxicity testing on the most sensitive aquatic organisms identified in the Phase IIA assessment, e.g.: the fish early life stage test, the Daphnia reproduction test, 72-hour algae tests and a bioaccumulation study,
- include, where an adequate safety margin between the PEC and PNEC values cannot be established, identification of effective mitigating measures to limit the environmental impact must be provided.

5. Section V: Form of monograph

5.1. Identity of the additive

5.1.1. Proposed proprietary name(s).

5.1.2. Type of additive according to its main function. Any other uses of the active substance should be specified.

5.1.3. Qualitative and quantitative composition (active substance, other components, impurities, batch to batch variation). If the active substance is a mixture of active components, each of which is clearly definable, the main components must be described separately and the proportions in the mixture given.

5.1.4. Physical state, particle size distribution, particle shape, density, bulk density; for liquids: viscosity, surface tension.

5.1.5. Manufacturing process including any specific processing procedures.

5.2. Specifications concerning the active substance

5.2.1. Generic name, chemical name according to IUPAC nomenclature, other generic international names and abbreviations. Chemical Abstracts Service Number (CAS).

5.2.2. Structural formula, molecular formula and molecular weight. Qualitative and quantitative composition of the main components, microbial origin (name and place of culture collection where the strain is deposited), if the active substance is a fermentation product.

5.2.3. Purity

Qualitative and quantitative composition of the active substances and occurring accompanying impurities and toxic substances, confirmation of the absence of the production organisms.

5.2.4. Relevant properties

Physical properties of the chemically specified substances: dissociation constant, pKa, electrostatic properties, melting point, boiling point, density, vapour pressure, solubility in water and organic solvents, K_{ow} and K_{oc} , mass and absorption spectra, NMR data, possible isomers and any other appropriate physical properties.

5.3. *Physico-chemical, technological and biological properties of the additive*

5.3.1. Stability of the additive on exposure to environmental conditions such as light, temperature, pH, moisture and oxygen. Proposal of a shelf life.

5.3.2. Stability during the preparation of premixtures and feedingstuffs, in particular stability to anticipated process conditions (heat, moisture, pressure/shear and time). Possible degradation or decomposition products.

5.3.3. Stability during the storage of premixtures and processed feedingstuffs under defined conditions. Proposal of a shelf life.

5.3.4. Other appropriate physico-chemical, technological or biological properties such as dispersability under favourable conditions in order to obtain and keep homogeneous mixtures in premixtures and feedingstuffs, antidusting and antistatic properties, dispersability in liquids.

5.4. *Control methods*

5.4.1. Description of the methods used for the determination of the criteria listed under items 2.1.3, 2.1.4, 2.2.3, 2.2.4, 2.3.1, 2.3.2, 2.3.3 and 2.3.4.

5.4.2. Description of the qualitative and quantitative analytical methods for determining the marker residue of the active substance in target tissues and animal produce.

5.4.3. If the said methods have been published the literature references may suffice and the corresponding reprints should be given.

5.4.4. Information on the optimum storage conditions for the reference standards.

5.5. *Biological properties of the additive*

5.5.1. Particulars of the prophylactic effects for coccidiostats and other medicinal substances (e.g. morbidity, mortality, oocyst count and lesion score).

5.5.2. For zootechnical additives other than those listed in 5.5.1 particulars of the effects on feed intake, body weight, feed efficiency, product quality and yield and any other parameter of benefit to the animal, the environment, the producer or the consumer.

5.5.3. For technological additives, relevant technological effects.

5.5.4. Any adverse effects, contra-indications or warnings (target animal, consumer, environment), including biological interactions, with particulars of their justification. Any ADI or MRLs established for other uses of the active substance should be specified.

5.6. *Details of the quantitative and qualitative residues in target tissues, if any, found in animal produce following envisaged use of the additive*

5.7. *The ADI, the established MRLs and the withdrawal period should be given, if appropriate*

5.8. *Other characteristics suitable for identification of the additive*

5.9. *Conditions of use*

5.10. *Date*

6. **Section VI: Form of identification note**

1. *Identity of the additive*

1.1. Type of additive

1.2. Physical state

1.3. Qualitative and quantitative composition

1.4. Method of analysis of the additive and the residues

1.5. Community registration number (EC number)

1.6. Packaging

2. *Specifications concerning the active substance*

2.1. Generic name, chemical name, CAS Number

— Generic name

— Chemical name (IUPAC)

— CAS number

2.2. Empirical formula

3. *Physico-chemical, technological and biological properties of the additive*

3.1. Stability of additive

3.2. Stability during the preparation of premixtures and feedingstuffs

3.3. Stability during storage of premixtures and feedingstuffs

3.4. Other properties

4. *Conditions of use*

4.1. Species or category of animals, maximum age if specified

4.2. Minimum and maximum content in feedingstuffs

4.3. Contra-indications, interactions

4.4. Warnings

5. *Person responsible for putting into circulation*

5.1. Name

5.2. Address

5.3. Registration number

6. *Manufacturer*

6.1. Name

6.2. Address

6.3. Approval number or registration number assigned to the establishment or the intermediary.

7. *Date*

7. **Section VII: Renewal of authorisation of additives whose authorisation is linked to a person responsible for putting them into circulation**

1. *General*

An updated dossier and monograph should be prepared according to the most up-to-date guidelines and a list provided of all variations of any type since the authorisation for putting into circulation or the last renewal.

It must be confirmed that the monograph and safety file has have been adapted to include all new information relevant to the additive or now required as a result of changes in these guidelines.

Information must also be provided on the authorisation status world wide and sales volume.

2. *Identity of the active substance and of the additive*

Evidence should be presented to show that the additive has not been changed or altered in composition, purity or activity in respect of the additive authorised. Any change of the manufacturing process should be reported.

3. *Efficacy*

Evidence should be presented to show that the additive retains the claimed efficacy under conditions of animal production current in the European Union at the time of application for renewal of the authorisation. This should include an account of general experience with the use of the additive and performance monitoring.

4. *Microbiology*

Special regard should be given to possible development of resistance to antimicrobials during the long-term use under practical conditions. The tests must thus be performed under field conditions in farms, where the additive has been routinely used for as long a time as possible. A selection of common intestinal bacteria should be used as test organisms, and the selection should include relevant endogenous and exogenous gram-positive as well as gram-negative organisms.

If the tests show a change in the resistance pattern compared to the original figures, the resistant bacteria must be examined for cross-resistance to relevant antibiotics used for treatment of infectious diseases in man and animals. The most important are antibiotics belonging to the same group as the additive, but also other groups of antibiotics should be included in the trial.

Results of appropriate monitoring programmes should be reported.

5. *Safety*

Evidence should be presented that in the light of the current knowledge the additive remains safe under the approved conditions for target species, consumers, operators and the environment. A safety update for the period since the authorisation for putting into circulation or the last renewal with information on the following items should be presented:

- reports on adverse effects including accidents (previously unknown effects, severe effects of any type, increased incidence of known effects) for target animals, operators and the environment. The report on adverse effect should include the nature of the effect, number of affected individuals/organisms, outcome, conditions of use, causality assessment;
- reports on previously unknown interactions and cross-contaminations;
- data from residue monitoring where appropriate;
- any other information concerning the safety of the additive.

If no further information is provided on any of these factors, the reasons for this should be clearly identified.

8. **Section VIII: New applicant relying on the first authorisation of an additive whose authorisation is linked to a person responsible for putting them into circulation**

Since reliance can be placed on the evaluation of the data supplied for initial authorisation a dossier prepared in relation to an application under Article 9c(3) need comply only with the following requirements.

An additive can be considered as identical for this purpose if the qualitative and quantitative composition and the purity of active and inactive components are essentially similar, the preparation is the same and the conditions of use are identical.

For such products it will normally not be necessary to repeat pharmacological, toxicological and efficacy studies and an abridged application can be submitted. This must include expert reports.

- A complete Section II and a monograph must be submitted.
- Data must be provided indicating that the specification range of the physical, chemical characteristics of the additive is essentially similar to that of the established product.
- It must be confirmed that further scientific knowledge in the available literature on the additive has not changed the original assessment on efficacy since the authorisation for putting into circulation of the original additive.
- Special regard should be given to possible development of resistance to antimicrobials during the long-term use of the active substance under practical conditions. The tests must thus be performed under field conditions in farms, where the active substance has been routinely used for as long a time as possible. A selection of common intestinal bacteria should be used as test organisms, and the selection should include relevant endogenous and exogenous gram-positive as well as gram-negative organisms.
- If the tests show a change in the resistance pattern compared to the original figures, the resistant bacteria must be examined for cross-resistance to relevant antibiotics used for treatment of infectious diseases in man and animals. The most important are antibiotics belonging to the same group as the additive, but also other groups of antibiotics should be included in the trial.
- Evidence should be presented that in the light of the current scientific knowledge in the available literature the additive remains safe under the approved conditions for target species, consumers, operators and the environment.
- The conformity of the withdrawal period with the MRL has to be established.

PART II

MICRO-ORGANISMS AND ENZYMES⁽¹⁾

⁽¹⁾ See Commission Directive 94/40/EC (OJ L 208, 11.8.1994, p. 15), as amended by Directive 95/11/EC.