Commission Regulation (EC) No 1451/2007 of 4 December 2007 on the second phase of the 10-year work programme referred to in Article 16(2) of Directive 98/8/EC of the European Parliament and of the Council concerning the placing of biocidal products on the market (Text with EEA relevance) (repealed)

Article 1	Subject matter
Article 2	Definitions
Article 3	Existing active substances
Article 3a	Procedure for the declarati

- Article 3a Procedure for the declaration of intention to notify
- Article 3b Notification procedure
- Article 3c Inclusion in, or exclusion from, the review programme
- Article 4 Non-inclusion
- Article 5 Derogation for essential use
- Article 6 Food and Feed
- Article 7 Examination of existing active substances under the review programme
- Article 8 Preparation of the complete dossier
- Article 9 Submission of the complete dossier
- Article 10 Joining and replacing of participants
- Article 11 Withdrawal of participants
- Article 12 Taking over the role of participant
- Article 13 Completeness check of dossiers
- Article 14 Evaluation of dossiers by the Rapporteur Member State
- Article 15 Commission procedures
- Article 16 Access to information
- Article 17 Suspension of procedures
- Article 18 Repeal
- Article 19 Entry into force
 - Signature

ANNEX I

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ANNEX II

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ANNEX III

Requirements for the complete dossier and the summary dossier

(a)

- (b) The summary dossier must include the following:
- (c)
- (d)

There are currently no known outstanding effects for the Commission Regulation (EC) No 1451/2007 (repealed).