Commission Regulation (EC) No 2032/2003 of 4 November 2003 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the placing of biocidal products on the market, and amending Regulation (EC) No 1896/2000 (Text with EEA relevance) (repealed)

Article 1	Subject matter				
Article 2	Definitions				
Article 3	Existing Active Substances				
Article 4	Non-inclusion				
Article 4a	Essential use				
Article 4b	Extension of deadline for placing on the market				
Article 5	Review of notified existing active substances				
Article 6	Preparation of the complete dossier				
Article 7	Submission of the complete dossier				
Article 8	Joining, replacing or withdrawal of participants				
Article 9	Completeness check of dossiers				
Article 10	Evaluation of dossiers by the Rapporteur Member State				
Article 11	Commission procedures				
Article 12					
Article 13	Suspension of procedures				
Article 14	Amendments to Regulation (EC) No 1896/2000				
Article 15	Entry into force				
	Signature				
	ANNEX I				
	ANNEAI				
	EXISTING ACTIVE SUBSTANCES				
	ANNEX II				
	EXISTING ACTIVE SUBSTANCES AND PRODUCT TYPES INCLUDED IN THE REVIEW PROGRAMME				

ANNEX III

EXISTING ACTIVE SUBSTANCES THAT HAVE BEEN IDENTIFIED BUT IN RESPECT OF WHICH NO NOTIFICATION HAS BEEN ACCEPTED OR NO MEMBER STATE HAS INDICATED AN INTEREST AND EXISTING ACTIVE SUBSTANCES THAT HAD BEEN NOTIFIED BUT IN RESPECT OF WHICH NO DOSSIER WAS SUBMITTED OR ACCEPTED AS COMPLETE AND FOR WHICH NO PRODUCER,

SPAIN

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EC) No 2032/2003 (repealed). (See end of Document for details)

FORMULATOR,	ASSOCIATION,	OTHER PE	ERSON OR	MEMBER	STATE HAS
INDICATED A	N INTEREST TO	TAKE OVE	ER THE RO	LE OF PAI	RTICIPANT

	INDICATED AN INTEREST TO TAKE OVER THE ROLE OF PARTICIPANT
	ANNEX IV
RE	QUIREMENTS FOR THE COMPLETE DOSSIER AND THE SUMMARY DOSSIER
(a)	
(b)	The summary dossier must include the following:
(c)	
(d)	
	ANNEX V
	TIME PERIODS AND RAPPORTEUR MEMBER STATES FOR THE SUBMISSION OF COMPLETE DOSSIERS FOR EXISTING ACTIVE SUBSTANCES INCLUDED IN THE REVIEW PROGRAMME
	Part A
	Part B
	Part C
	Part D
	ANNEX VI
	COMPETENT AUTHORITIES, AS REFERRED TO IN ARTICLE 5(4) AS OF 1 MARCH 2005
	BELGIUM
	CZECH REPUBLIC
	DENMARK
	GERMANY
	ESTONIA
	GREECE

Document Generated: 2023-12-27

Changes to legislation: There are currently no known outstanding effects for the

Commission Regulation (EC) No 2032/2003 (repealed). (See end of Document for details)

FRANCE IRELAND ITALY CYPRUS LATVIA LITUANIA LUXEMBOURG **HUNGARY MALTA NETHERLANDS AUSTRIA POLAND PORTUGAL SLOVENIA SLOVAKIA FINLAND SWEDEN** UNITED KINGDOM **ICELAND NORWAY** ANNEX VII EXISTING ACTIVE SUBSTANCES THAT WERE NOT IDENTIFIED BY 28 MARCH 2002 BUT MAY REMAIN ON THE MARKET UNTIL 1 SEPTEMBER 2006 ANNEX VIII

EXISTING ACTIVE SUBSTANCES THAT WERE NOTIFIED BUT IN RESPECT OF WHICH NO DOSSIER WAS SUBMITTED OR ACCEPTED AS COMPLETE AND FOR WHICH ANOTHER PRODUCER, FORMULATOR,

Changes to legislation: There are currently no known outstanding effects for the Commission Regulation (EC) No 2032/2003 (repealed). (See end of Document for details)

ASSOCIATION, OTHER PERSON OR MEMBER STATE HAS INDICATED AN INTEREST TO TAKE OVER THE ROLE OF PARTICIPANT

Changes to legislation:

There are currently no known outstanding effects for the Commission Regulation (EC) No 2032/2003 (repealed).