Commission Regulation (EU) No 545/2011 of 10 June 2011 implementing Regulation (EC) No 1107/2009 of the European Parliament and of the Council as regards the data requirements for plant protection products (Text with EEA relevance) (repealed)

Article 1 Article 2	Signature
	ANNEX
	A REQUIREMENTS FOR PLANT PROTECTION PRODUCTS, AS DED FOR IN ARTICLE 8(1)(c) OF REGULATION (EC) No 1107/2009
	DDUCTION
1. 2.	Tests and analyses
2.	2.1
	<ul> <li>2.2. Tests and analyses, required under points 6.2 to 6.7 of</li> <li>2.3. Officially recognised testing facilities and organisations, and, where requested, official</li> </ul>
	2.4
	2.5
	2.6
3.	
4.	
	PART A
	CHEMICAL PREPARATIONS
Identit	y of the plant protection product
1.1.	Applicant (name and address, etc.)
1.1.	Manufacturer of the preparation and the active substance(s) (names and
1.3.	Trade name or proposed trade name, and manufacturer's development code
1.4.	Detailed quantitative and qualitative information on the composition of the
	1.4.1
	1.4.2
	1.4.3
	1.4.4
1.5.	Physical state and nature of the preparation (emulsifiable concentrate, wettable 1.5.1
1.6.	Function (herbicide, insecticide, etc.)
Physic	al, chemical and technical properties of the plant protection product
2.1.	Appearance (colour and odour)
2.2.	Explosivity and oxidising properties
	2.2.1

1.

2.

3.

Document Generated: 2023-11-25

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

		2.2.2.		
2.3	3.		point and other indications of flammability or spontaneous ignition	
	4.		y/alkalinity and if necessary pH value	
2.5	5.		ity and surface tension	
26	5.	Relativ	ve density and bulk density	
	•			
2.7	7		e — stability and shelf-life: Effects of light, temperature and	
2.,				
2.8	2		cal characteristics of the plant protection product	
2.0	٥.		Wettability	
			Persistent foaming	
			Suspensibility and suspension stability	
			Dilution stability	
			Dry sieve test and wet sieve test	
			Particle size distribution (dustable and wettable powders, granules),	
		2.8.0.	content of	
			2.8.6.1	
			2.8.6.2	
		207	2.8.6.3	
		2.8.7.	Emulsifiability, re-emulsifiability, emulsion stability	
			2.8.7.1	
		200	2.8.7.2	
		2.8.8.	Flowability, pourability (rinsability) and dustability	
			2.8.8.1	
			2.8.8.2	
2.0	`	DI '	2.8.8.3	
2.9	1.	•	al and chemical compatibility with other products including plant	
		protect		
2 1		2.9.2.	1.11 - 11 - 1	
2.1			ence and distribution to seeds	
2.1	H.	Summ	ary and evaluation of data presented under points 2.1 to	
ъ		1.		
		n applica		
3.1			of use envisaged, e.g. field, protected crops, storage of	
3.2			s on harmful organisms, e.g. contact, inhalation or stomach poison,	
	Details of intended use, e.g. types of harmful organisms controlled  Application rate			
3.5				
3.6			d of application	
3.7			er and timing of applications and duration of protection	
3.8			sary waiting periods or other precautions to avoid phytotoxic effects	
3.9	€.	Propos	ed instructions for use	

4. Further information on the plant protection product

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

	4.1.	Packaging (type, materials, size, etc.), compatibility of the preparation with 4.1.1. 4.1.2. 4.1.2.
	4.0	4.1.3
	4.2.	Procedures for cleaning application equipment
	4.3.	Re-entry periods, necessary waiting periods or other precautions to protect 4.3.1
	4.4.	4.3.2
	4.5.	Emergency measures in the case of an accident
	4.6.	Procedures for destruction or decontamination of the plant protection product.  4.6.1. Possibility of neutralisation
		<ul><li>4.6.2. Controlled incineration</li><li>4.6.3. Others</li></ul>
5.	Analy	tical methods
		Introduction
	5.1.	Methods for the analysis of the preparation 5.1.1.
		5.1.2
		5.1.3. Specificity, linearity, accuracy and repeatability 5.1.3.1
		5.1.3.2
		5.1.3.4
	5.2.	Analytical methods for the determination of residues
6.	Effica	cy data
		General
	6.1.	J
	6.2.	$\boldsymbol{\mathcal{C}}$
		Aim of the tests
		Test conditions
		Test guideline
	6.3.	Information on the occurrence or possible occurrence of the development
	6.4.	Effects on the yield of treated plants or plant products
		6.4.1. Effects on the quality of plants or plant products Aim of the tests
		Circumstances in which required
		6.4.2. Effects on transformation processes
		Aim of the tests
		Circumstances in which required
		6.4.3. Effects on the yield of treated plants or plant products  Aim of the tests
		Circumstances in which required
	6.5.	Phytotoxicity to target plants (including different cultivars), or to target
		Aim of the tests
		Circumstances in which required
		Test guideline
	6.6.	Observations on undesirable or unintended side-effects, e.g. on beneficial and.
		6.6.1. Impact on succeeding crops  Aim of the information required

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

Circumstances in which required

6.6.2. Impact on other plants, including adjacent crops

Aim of the information required

Circumstances in which required

6.6.3. Impact on treated plants or plant products to be used...

Aim of the information required Circumstances in which required Test guideline

6.6.4. Effects on beneficial and other non-target organisms

6.7. Summary and evaluation of data presented under 6.1 to 6.6...

### 7. Toxicological studies

## 7.1. Acute toxicity

7.1.1. Oral

Circumstances in which required

Test method

7.1.2. Percutaneous

Circumstances in which required

Test method

7.1.3. Inhalation

Aim of the test

Circumstances in which required

Test method

7.1.4. Skin irritation

Aim of the test

Circumstances in which required

Test method

7.1.5. Eye irritation

Aim of the test

Circumstances in which required

Test method

7.1.6. Skin sensitisation

Aim of the test

Circumstances in which required

Test method

7.1.7. Supplementary studies for combinations of plant protection products

Aim of the test

## 7.2. Data on exposure

7.2.1. Operator exposure

7.2.1.1. Estimation of operator exposure

Aim of the estimation

Circumstances in which required

**Estimation conditions** 

7.2.1.2. Measurement of operator exposure

Aim of the test

Circumstances in which required

Test conditions

7.2.2. Bystander exposure

Aim of the estimation

Circumstances in which required

**Estimation conditions** 

7.2.3. Worker exposure

7.2.3.1. Estimation of worker exposure

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

Aim of the estimation

Circumstances in which required

**Estimation conditions** 

7.2.3.2. Measurement of worker exposure

Aim of the test

Circumstances in which required

Test conditions

7.3. Dermal absorption

Aim of the test

Circumstances in which required

Test conditions

Test guideline

- 7.4. Available toxicological data relating to non-active substances
- 8. Residues in or on treated products, food and feed

Introduction

8.1. Metabolism, distribution and expression of residue in plants or livestock...

Aim of the tests

Circumstances in which required

Test conditions

8.2. Residue trials

Aim of the tests

Circumstances in which required

Test conditions

8.3. Livestock feeding studies

Aim of the tests

Circumstances in which required

Test conditions

8.4. Effects of industrial processing and/or household preparations

Aim of the tests

Circumstances in which required

Test conditions

8.5. Residues in succeeding crops

Aim of the test

Circumstances in which required

Test conditions

- 8.6. Proposed maximum residue levels (MRLs) and residue definition
- 8.7. Proposed pre-harvest intervals for envisaged uses, or withholding periods or...
- 8.8. Estimation of the potential and actual exposure through diet and...
- 8.9. Summary and evaluation of residue behaviour
- 9. Fate and behaviour in the environment

Introduction

- (i) ......
- (ii) In particular, the information provided for the plant protection product,...
- (iv) Where relevant tests shall be designed and data analysed using...
- (v) Predicted environmental concentrations in soil (PECS), water (PECSW and PECGW)...
- 9.1. Fate and behaviour in soil
  - 9.1.1. Rate of degradation in soil
    - 9.1.1.1. Laboratory studies

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

Aim of the test Circumstances in which required Test conditions Test guideline 9.1.1.2. Field studies Soil dissipation studies Aim of the test Circumstances in which required Test conditions and test guideline Soil residue studies Aim of the test Circumstances in which required Test conditions Test guideline Soil accumulation studies Aim of the tests Circumstances in which required Test conditions Test guideline 9.1.2. Mobility in the soil Aim of the test 9.1.2.1. Laboratory studies Circumstances in which required Test guideline 9.1.2.2. Lysimeter studies or field leaching studies Aim of the tests Circumstances in which required Test conditions Estimation of expected concentrations in soil 9.2. Fate and behaviour in water 9.2.1. Estimation of concentrations in groundwater 9.2.2. Impact on water treatment procedures 9.2.3. Estimation of concentrations in surface water 9.3. Fate and behaviour in air 10. Ecotoxicological studies Introduction (i) In particular, the information provided for the plant protection (ii) product,... (iii) In general, much of the data relating to impact on... (iv) (v) (vi) Where exposure data are necessary to decide whether a study... (vii) (viii) (ix) 10.1. Effects on birds 10.1.1. Acute oral toxicity Aim of the test Circumstances in which required Test conditions 10.1.2. Supervised cage or field trials

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

Aim of the test

Circumstances in which required

Test conditions

10.1.3. Acceptance of bait, granules or treated seeds by birds

Aim of the test

Circumstances in which required

10.1.4. Effects of secondary poisoning

10.2. Effects on aquatic organisms

10.2.1. Acute toxicity to fish, aquatic invertebrates or effects on algal...

Circumstances in which required

Test conditions and test guidelines

10.2.2. Microcosm or mesocosm study

Aim of the test

Circumstances in which required

Test conditions

Test guideline

10.2.3. Residue data in fish

Aim of the test

Circumstances in which required

Test guideline

10.2.4. Additional studies

10.3. Effects on terrestrial vertebrates other than birds

Aim of the test

Circumstances in which required

Test conditions

10.4. Effects on bees

10.4.1. Acute oral and contact toxicity

Aim of the test

Circumstances in which required

Test guideline

10.4.2. Residue test

Aim of the test

Circumstances in which required

Test conditions

10.4.3. Cage tests

Aim of the test

Circumstances in which required

Test conditions

Test guideline

10.4.4. Field tests

Aim of the test

Circumstances in which required

Test conditions

Test guideline

10.4.5. Tunnel tests

Aim of the test

Circumstances in which required

Test conditions

Test guideline

10.5. Effects on arthropods other than bees

10.5.1. Laboratory, extended laboratory and semi-field tests

Aim of the test

Circumstances in which required

(iv)

(v)

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

Test conditions Test guideline 10.5.2. Field tests Aim of the test Circumstances in which required Test conditions Test guideline Effects on earthworms and other soil non-target macro-organisms, believed to... 10.6. 10.6.1. Effects on earthworms 10.6.1.1 Acute toxicity tests Aim of the test Circumstances in which required Test guideline 10.6.1.2Tests for sublethal effects Aim of the test Circumstances in which required Test conditions 10.6.1.3Field studies Aim of the test Circumstances in which required Test conditions 10.6.2. Effects on other soil non-target macro-organisms Aim of the test Circumstances in which required 10.7. Effects on soil non-target micro-organisms 10.7.1. Laboratory testing Aim of the test Circumstances in which required Test guideline 10.7.2. Additional testing Aim of the test Circumstances in which required Available data from biological primary screening in summary form 10.8. 11. Summary and evaluation of Sections 9 and 10 Further information 12. Information on authorisations in other countries 12.1 Information on established maximum residue levels (MRL) in other countries... 12.2. 12.3. Proposals including justification for the classification and labelling proposed in... 12.4. 12.5. Specimens of proposed packaging PART B PREPARATIONS OF MICRO-ORGANISMS INCLUDING VIRUSES Introduction This Part provides data requirements for the authorisation of a... (i) (ii) (iii) 

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

	(vi)	
1.	1.1. 1.2. 1.3. 1.4.	Applicant Manufacturer of the preparation and the micro-organism(s) Trade name or proposed trade name, and manufacturer's development code Detailed quantitative and qualitative information on the composition of the (i) (ii) For preparations the following information must be reported: (iii)
	1.6.	Function
2.	Physica 2.1. 2.2. 2.3. 2.4. 2.5. 2.6. 2.7.	Appearance (colour and odour) Storage stability and shelf-life 2.2.1. Effects of light, temperature and humidity on technical characteristics of  (i)
		(i) The size distribution of particles in the case of powders, (ii)
	2.8.	Physical, chemical and biological compatibility with other products including plant  2.8.1. Physical compatibility  2.8.2. Chemical compatibility  2.8.3. Biological compatibility
	2.9.	Adherence and distribution to seeds
	2.10.	Summary and evaluation of data presented under points 2.1 to

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

3.	Data	οn	ann	licatio	n
<b>)</b> .	Data	on	app.	псано	n

- 3.1. Field of use envisaged
- 3.2. Mode of action
- 3.3. Details of intended use
- 3.4. Application rate
- 3.5. Content of micro-organism in material used (e.g. in the diluted...
- 3.6. Method of application
- 3.7. Number and timing of applications and duration of protection
- 3.8. Necessary waiting periods or other precautions to avoid phytopathogenic effects...
- 3.9. Proposed instructions for use

## 4. Further information on the plant protection product

- 4.1. Packaging and compatibility of the preparation with proposed packaging materials...

  - (iii) .....
- 4.2. Procedures for cleaning application equipment
- 4.3. Re-entry periods, necessary waiting periods or other precautions to protect...
- 4.4. Recommended methods and precautions concerning: handling, storage, transport or fire...
- 4.5. Measures in the case of an accident
- 4.6. Procedures for destruction or decontamination of the plant protection product...
  - 4.6.1. Controlled incineration
  - 4.6.2. Others
- 5. Analytical methods

Introduction

- 5.1. Methods for the analysis of the preparation
- 5.2. Methods to determine and quantify residues
- 6. Efficacy data

General

- 6.1. Preliminary tests
- 6.2. Testing effectiveness

Aim of the tests

Test conditions

Test guideline

- 6.3. Information on the occurrence or possible occurrence of the development...
- 6.4. Effects on the yield of treated plants or plant products...
  - 6.4.1. Effects on the quality of plants or plant products

Aim of the tests

Circumstances in which required

6.4.2. Effects on transformation processes

Aim of the tests

Circumstances in which required

6.4.3. Effects on the yield of treated plants or plant products...

Aim of the tests

Circumstances in which required

6.5. Phytotoxicity to target plants (including different cultivars), or to target...

Aim of the tests

Circumstances in which required

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

#### Test guideline

- 6.6. Observations on undesirable or unintended side-effects, e.g. on beneficial and...
  - 6.6.1. Impact on succeeding crops

Aim of the information required

Circumstances in which required

6.6.2. Impact on other plants, including adjacent crops

Aim of the information required

Circumstances in which required

6.6.3. Impact on treated plants or plant products to be used...

Aim of the information required

Circumstances in which required

Test guideline

- 6.6.4. Effects on beneficial and other non-target organisms
- 6.7. Summary and evaluation of data presented under 6.1 to 6.6...
- 7. Effects on human health
  - 7.1. Basic acute toxicity studies
    - 7.1.1. Acute oral toxicity

Circumstances in which required

Test method

7.1.2. Acute inhalation toxicity

Aim of the test

Circumstances in which required

Test method

7.1.3. Acute percutaneous toxicity

Circumstances in which required

Test method

- 7.2. Additional acute toxicity studies
  - 7.2.1. Skin irritation

Aim of the test

Circumstances in which required

Test method

7.2.2. Eye irritation

Aim of the test

Circumstances in which required

Test method

7.2.3. Skin sensitisation

Aim of the test

Circumstances in which required

Test method

- 7.3. Data on exposure
- 7.4. Available toxicological data relating to non-active substances
- 7.5. Supplementary studies for combinations of plant protection products Aim of the test
- 7.6. Summary and evaluation of health effects
- 8. Residues in or on treated products, food and feed
- 9. Fate and behaviour in the environment
- 10. Effects on non-target organisms

Introduction

(i) .....

11.

Document Generated: 2023-11-25

Changes to legislation: There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

	<ul><li>(ii)</li></ul>
	(iv)
	(v)
	(vi) Where exposure data are necessary to decide whether a study
	(vii)
10.1.	Effects on birds
10.2.	Effects on aquatic organisms
10.3.	Effects on bees
10.4.	Effects on arthropods other than bees
10.5.	Effects on earthworms
10.6.	Effects on soil micro-organisms
10.7.	Additional studies

Summary and evaluation of environmental impact

#### **Changes to legislation:**

There are outstanding changes not yet made to Commission Regulation (EU) No 545/2011 (repealed). Any changes that have already been made to the legislation appear in the content and are referenced with annotations.

View outstanding changes

## Changes and effects yet to be applied to:

- Annex word inserted by S.I. 2019/556 reg. 17(6)(g)(ii)
- Annex Pt. A point 1.4.3 word substituted by S.I. 2019/720 Sch. 2 para. 59(2)(b)(ii)
- Annex Pt. B point 1.4(iii) word substituted by S.I. 2019/720 Sch. 2 para. 59(2)(c)(ii)
- Annex word substituted in earlier amending provision S.I. 2019/720, Sch. 2 para.
   59(2)(a) by S.I. 2020/1567 Sch. 2 para.
- Annex Pt. A point 1.4.3 word substituted in earlier amending provision S.I. 2019/720, Sch. 2 para. 59(2)(b)(i) by S.I. 2020/1567 Sch. 2 para. 19
- Annex Pt. B point 1.4.4(iii) word substituted in earlier amending provision S.I. 2019/720, Sch. 2 para. 59(2)(c)(i) by S.I. 2020/1567 Sch. 2 para. 19
- Annex words inserted by S.I. 2019/556 reg. 17(6)(d)
- Annex words inserted by S.I. 2019/556 reg. 17(6)(f)(ii)(bb)
- Annex Pt. A point 7.2 words inserted by S.I. 2019/556 reg. 17(7)(e)
- Annex Pt. A point 7.2.1.2 words inserted by S.I. 2019/556 reg. 17(7)(g)
- Annex Pt. A point 7.2.3.2 words inserted by S.I. 2019/556 reg. 17(7)(g)
- Annex Pt. A point 7.3 words inserted by S.I. 2019/556 reg. 17(7)(h)
- Annex words inserted by S.I. 2019/720 Sch. 2 para. 59(2)(a)
- Annex words omitted by S.I. 2019/556 reg. 17(6)(h)
- Annex Pt. A point 7.1 words omitted by S.I. 2019/556 reg. 17(7)(d)
- Annex Pt. A point 7.1.1 words omitted by S.I. 2019/556 reg. 17(7)(d)
- Annex Pt. A point 7.1.2 words omitted by S.I. 2019/556 reg. 17(7)(d)
- Annex Pt. A point 7.2.1.1 words omitted by S.I. 2019/556 reg. 17(7)(f)
- Annex Pt. A point 8.9 words omitted by S.I. 2019/556 reg. 17(7)(i)
- Annex Pt. A point 12.3 words omitted by S.I. 2019/556 reg. 17(7)(k)
- Annex Pt. B point 7.1 words omitted by S.I. 2019/556 reg. 17(8)(d)
- Annex Pt. B point 7.1.1 words omitted by S.I. 2019/556 reg. 17(8)(d)
- Annex Pt. B point 7.1.3 words omitted by S.I. 2019/556 reg. 17(8)(d)
- Annex words substituted by S.I. 2019/556 reg. 17(6)(a)
- Annex words substituted by S.I. 2019/556 reg. 17(6)(b)
- Annex words substituted by S.I. 2019/556 reg. 17(6)(c)
- Annex words substituted by S.I. 2019/556 reg. 17(6)(e) (This amendment not applied to legislation.gov.uk. Reg. 17(6)(e) substituted immediately before IP completion day by S.I. 2020/1376, regs. 1(4), 3(15)(b))
- Annex words substituted by S.I. 2019/556 reg. 17(6)(f)(i)
- Annex words substituted by S.I. 2019/556 reg. 17(6)(f)(ii)(aa)
- Annex words substituted by S.I. 2019/556 reg. 17(6)(g)(i)
- Annex words substituted by S.I. 2019/556 reg. 17(6)(i)
- Annex Pt. A point 1.1 words substituted by S.I. 2019/556 reg. 17(7)(a)
- Annex Pt. A point 1.4.1 words substituted by S.I. 2019/556 reg. 17(7)(b)
- Annex Pt. A point 6.5 words substituted by S.I. 2019/556 reg. 17(7)(c)(i)
- Annex Pt. A point 6.5 words substituted by S.I. 2019/556 reg. 17(7)(c)(ii)
- Annex Pt. B point 1.1 words substituted by S.I. 2019/556 reg. 17(8)(a)
- Annex Pt. B point 1.4(ii) words substituted by S.I. 2019/556 reg. 17(8)(b)
- Annex Pt. B point 6.5 words substituted by S.I. 2019/556 reg. 17(8)(c)(i)
- Annex Pt. B point 6.5 words substituted by S.I. 2019/556 reg. 17(8)(c)(ii)
- Annex Pt. A point 1.4.3 words substituted by S.I. 2019/720 Sch. 2 para. 59(2)(b)(i)
- Annex Pt. B point 1.4(iii) words substituted by S.I. 2019/720 Sch. 2 para. 59(2)(c)(i)
- Annex words substituted by S.I. 2019/556, reg. 17(6)(e) (as substituted) by S.I. 2020/1376 reg. 3(15)(b)
- Annex Pt. A point 1.1 words substituted in earlier amending provision S.I. 2019/556, reg. 17(7)(a) by S.I. 2020/1376 reg. 3(15)(c)(i)

- Annex Pt. A point 1.1 words substituted in earlier amending provision S.I. 2019/556, reg. 17(7)(a) by S.I. 2020/1376 reg. 3(15)(c)(ii)
- Annex Pt. B point 1.1 words substituted in earlier amending provision S.I. 2019/556, reg. 17(8)(a) by S.I. 2020/1376 reg. 3(15)(c)(i)
- Annex Pt. B point 1.1 words substituted in earlier amending provision S.I. 2019/556, reg. 17(8)(a) by S.I. 2020/1376 reg. 3(15)(c)(ii)
- Art. 2 omitted by S.I. 2019/556 reg. 17(3)

# Changes and effects yet to be applied to the whole legislation item and associated provisions

- Signature words omitted by S.I. 2019/556 reg. 17(4)
- Art. A1 inserted by S.I. 2019/556 reg. 17(2)
- Art. A1 words substituted in earlier amending provision S.I. 2019/556, reg. 17(2) by S.I. 2020/1376 reg. 3(15)(a)
- Annex Pt. A s. 11 words omitted by S.I. 2019/556 reg. 17(7)(j)
- Annex Pt. B s. 11 words omitted by S.I. 2019/556 reg. 17(8)(e)