Information data sheet
according to article 32 of Regulation (EC) 1907/2006 (REACH)

Printing date 01.05.2026 Version number 3 (replaces version 2) Revision: 01.05.2026

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SECTION 1: Identification of the substance/mixture and of the company/undertaking

- **1.1 Product identifier**

- **Trade name: MUSKIL BR**

- **1.2 Relevant identified uses of the substance or mixture and uses advised against**
Ready for use rodenticide (biocidal product-PT14)

- **1.3 Details of the supplier of the information data sheet**

- **Manufacturer/Supplier:**
Zapi S.p.A.
Via Terza Strada, 12
35026 Conselve (PD) - Italy
Tel. +39 049 9597737 Fax +39 049 9597735

E-mail address of the competent person responsible for the information data sheet: techdept@zapi.it

- **Further information obtainable from:** Tech. dept.

- **1.4 Emergency telephone number:** Zapi customer service (Tel. +39 049 9597737): 9:00-12:00 / 14:00-17:00
Giftnotruf Berlin +49 (0) 30 30686700 Beratung in Deutsch und English.

SECTION 2: Hazards identification

- **2.1 Classification of the substance or mixture**

- **Classification according to Regulation (EC) No 1272/2008**
The product is not classified, according to the CLP regulation.

- **2.2 Label elements**

- **Labelling according to Regulation (EC) No 1272/2008** Not applicable

- **Hazard pictograms** Not applicable

- **Signal word** Not applicable

- **Hazard statements** Not applicable

- **2.3 Other hazards**

- **Results of PBT and vPvB assessment**

- **PBT:**	
56073-10-0 brodifacoum	
PBT	Brodifacoum fulfils the P, B and T criteria.

- **vPvB:**	
56073-10-0 brodifacoum	
vPvB	Brodifacoum fulfils the vP criterion.

- **Determination of endocrine-disrupting properties**
The mixture does not contain substances with endocrine disrupting properties in concentration equal to or greater than 0.1% by weight.

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SECTION 3: Composition/information on ingredients

- **3.2 Mixtures**

- **Description:** Mixture of substances listed below with nonhazardous additions.

- **Dangerous components:**		
CAS: 56073-10-0 EINECS: 259-980-5 Index number: 607-172-00-1	brodifacoum Acute Tox. 1, H300 (ATE=0.4 mg/kg bw); Acute Tox. 1, H310 (ATE=3.16 mg/kg bw); Acute Tox. 1, H330 (ATE=3.05 mg/m3); Repr. 1A, H360D; STOT RE 1, H372; Aquatic Acute 1, H400 (M=10); Aquatic Chronic 1, H410 (M=10); PBT, EUH440 Specific concentration limits: Repr. 1A; H360: C ≥ 0.003 % STOT RE 1; H372: C ≥ 0.02 % STOT RE 2; H373: 0.002 % ≤ C < 0.02 %	0.0017%

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- **Additional information:** For the wording of the listed hazard phrases refer to section 16.

SECTION 4: First aid measures

- 4.1 Description of first aid measures

- **General information:**
Please refer to the instructions below for each specific way of exposure.
If medical advice is needed, have product container or label at hand.

- **After inhalation:** Supply fresh air and to be sure call for a doctor.

- **After skin contact:**
Wash skin with water.
If symptoms occur call a POISON CENTRE or a doctor.

- **After eye contact:**
If symptoms occur rinse with water.
Remove contact lenses, if present and easy to do.
Call a POISON CENTRE or a doctor.

- **After swallowing:**
Rinse mouth.
If symptoms: Call 112/ambulance for medical assistance.
If no symptoms: call a POISON CENTRE or a doctor.
Information to Healthcare personnel/doctor: initiate life support measures if needed, thereafter call a POISON CENTRE.
Contact a veterinary surgeon in case of ingestion by a pet.

- 4.2 Most important symptoms and effects, both acute and delayed
This product contains an anticoagulant substance. If ingested, symptoms, which may be delayed, may include nosebleed and bleeding gums. In severe cases, there may be bruising and blood present in the faeces or urine.
Antidote: Vitamin K1 administered by medical/veterinary personnel only.

- 4.3 Indication of any immediate medical attention and special treatment needed
The primary treatment are the antidote therapy and the clinical assessment. Antidote: Vitamin K1 (phytomenadione). The effectiveness of the treatment should be monitored by measuring the clotting time. Do not interrupt the treatment until the clotting time is back to normality and is stable.
Consult a Poison Control Centre.

SECTION 5: Firefighting measures

- 5.1 Extinguishing media

- **Suitable extinguishing agents:** CO_2, powder or water spray. Fight larger fires with water spray.

- **For safety reasons unsuitable extinguishing agents:** To our knowledge, there are no unsuitable equipments.

- **5.2 Special hazards arising from the substance or mixture** In case of fire, toxic gases may be generated.

- **5.3 Advice for firefighters** Firefighters equipment in accordance with EN469 European standards.

- **Protective equipment:**
Do not inhale explosion gases or combustion gases.
Firefighters equipment in accordance with EN469 European standards.

- **Additional information**
Dispose of fire debris and contaminated fire fighting water in accordance with official regulations.

SECTION 6: Accidental release measures

- 6.1 Personal precautions, protective equipment and emergency procedures
Wear protective equipment. Keep unprotected persons away.

- 6.2 Environmental precautions:
Inform respective authorities in case of seepage into water course or sewage system.

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Do not allow to enter sewers/ surface or ground water.

- **6.3 Methods and material for containment and cleaning up:**
Pick up mechanically.
After cleaning up, ensure adequate ventilation.
Dispose of the material collected according to regulations.

- **6.4 Reference to other sections**
See Section 7 for information on safe handling.
See Section 8 for information on personal protection equipment.
See Section 13 for disposal information.

* SECTION 7: Handling and storage

- **7.1 Precautions for safe handling**
Do not smoke near the product.
When using the product, do not eat, drink or smoke.
Wash hands and directly exposed skin after using the product.
Place the product out of the reach of children, birds, pets and farm animals and other non-target animals.
Place the product away from food, drink and animal feeding stuffs, as well as from utensils or surfaces that have contact with these.
Do not use the product close to surface waters (e.g. rivers, ponds, water channels, dykes, irrigation ditches).
When placing the product close to water drainage systems, ensure that bait contact with water is avoided.

- **Information about fire - and explosion protection:**
See Section 6.
See section 5.

- **7.2 Conditions for safe storage, including any incompatibilities**

- **Requirements to be met by storerooms and receptacles:**
Store in a dry, cool and well ventilated place. Keep the container closed and away from direct sunlight.
Store in places prevented from the access of children, birds, pets and farm animals.

- **Information about storage in one common storage facility:**
Do not store near food, drink and feed.
Place the product away from food, drink and animal feeding stuffs, as well as from utensils or surfaces that have contact with these.

- **Further information about storage conditions:**
Protect from frost.
Protect from humidity and water.
Protect from light.
Store at temperatures not exceeding 35°C.

- **7.3 Specific end use(s)** This product is a rodenticide bait for rodents' control.

* SECTION 8: Exposure controls/personal protection

- **8.1 Control parameters**

- **Ingredients with limit values that require monitoring at the workplace:**
The product does not contain any substances for which there are Union workplace exposure limits.

- **PNECs**		
56073-10-0 brodifacoum		
Oral	PNEC	0.0000128 mg/kg bw (bird)
		0.000011 mg/kg bw (mammal)
	PNEC	0.00004 mg/l (aquatic organisms)
		>0.0038 mg/l (sewage treatment plant)
	PNEC	>0.88 mg/kg ww (soil)

- **Other exposure limit values**	
56073-10-0 brodifacoum	
AEL - long term	0.0000033 mg/kg bw/d

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AEL - medium term	0.0000067 mg/kg bw/d
AEL - short term	0.0000033 mg/kg bw/d

- **8.2 Exposure controls**

- **Appropriate engineering controls** No further data; see section 7.

- **Individual protection measures, such as personal protective equipment**

- **General protective and hygienic measures:**
The usual precautionary measures are to be adhered to when handling chemicals.
Keep away from food, drink and animal feedingstuffs.
Wash hands before breaks and at the end of work.
Do not eat, drink, smoke or sniff while working.

- **Respiratory protection:** Not required during normal use of the product.

- **Hand protection**

Professional use: wear protective chemical resistant gloves (EN 374) during product handling phase.

- **Material of gloves**
Glove materials: nitrile, polychloroprene (neoprene) or fluoroelastomer.
The thickness must, depending on the model and the type of fabric, generally be between 0.5 mm and 1.5 mm.

- **Penetration time of glove material**
The use of type B gloves, tested with minimum 3 substances (breakthrough time greater than 30 minutes according to EN 374) or higher is recommended.
WARNING:
When selecting specific gloves for use in particular applications and duration of use, other factors should be considered, such as (but not limited to): other chemicals handled, physical demands (cut/puncture protection, manual dexterity, thermal protection) possible body reactions to the glove material, as well as the instructions/specifications provided by the glove manufacturer.

- **Eye/face protection** Not required during normal use of the product.

- **Environmental exposure controls** See section 6.

- **Risk management measures** Follow the above-reported directions.

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SECTION 9: Physical and chemical properties

- **9.1 Information on basic physical and chemical properties**
- **General Information**

Physical state	Solid
Colour:	Light red
Odour:	Characteristic
Odour threshold:	No data available.
Melting point/freezing point:	No data available.
Boiling point or initial boiling point and boiling range	Not applicable (solid).
Flammability	Not flammable (EC 440/2008 No. A.10)
Lower and upper explosion limit	
Lower:	No data available.
Upper:	No data available.
Flash point:	Not applicable (solid).
Auto-ignition temperature:	Not applicable.
Decomposition temperature:	No data available.
pH at 20°C	6.6 (CIPAC MT 75.3 - 1% aq.)

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- **Viscosity:**	
- **Kinematic viscosity**	Not applicable.
- **Dynamic viscosity:**	Not applicable.
- **Solubility**	
- **water:**	Insoluble.
- **Partition coefficient n-octanol/water (log value)**	No data available.
- **Vapour pressure:**	Not applicable.
- **Density and/or relative density**	
- **Density at 20°C:**	1.440 g/mL (EC 440/2008 No. A.3).
- **Relative density**	No data available.
- **Vapour density**	Not applicable.
- **Particle characteristics**	No data available
- **9.2 Other information**	
- **Appearance:**	
- **Form:**	Solid
- **Information with regard to physical hazard classes**	
- **Explosives**	Not explosive
- **Flammable gases**	Not applicable
- **Aerosols**	Not applicable
- **Oxidising gases**	Not applicable
- **Gases under pressure**	Not applicable
- **Flammable liquids**	Not applicable
- **Flammable solids**	Not flammable.
- **Self-reactive substances and mixtures**	Not self-reactive
- **Pyrophoric liquids**	Not applicable
- **Pyrophoric solids**	Not pyrophoric
- **Self-heating substances and mixtures**	Not self-heating
- **Substances and mixtures, which emit flammable gases in contact with water**	Not applicable
- **Oxidising liquids**	Not applicable
- **Oxidising solids**	Not oxidising
- **Organic peroxides**	Not applicable
- **Corrosive to metals**	Not applicable
- **Desensitised explosives**	Not applicable

SECTION 10: Stability and reactivity

- **10.1 Reactivity** Under standard handling and storing conditions, the product does not show any dangerous reaction.
- **10.2 Chemical stability** Stable at room temperature and if used as recommended.
- **Thermal decomposition / conditions to be avoided:** No decomposition if used according to specifications.
- **10.3 Possibility of hazardous reactions** No dangerous reactions known.
- **10.4 Conditions to avoid**
  Under standard handling and storing conditions, the product does not show any dangerous reaction.
- **10.5 Incompatible materials:**
  Store only in original container.
  Given the lack of information about possible incompatibilities with other substances, it is recommended not to use it in combination with other products.
- **10.6 Hazardous decomposition products:**
  No dangerous decomposition products known under normal conditions of storage and use.

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SECTION 11: Toxicological information

- 11.1 Information on hazard classes as defined in Regulation (EC) No 1272/2008

- **Acute toxicity** Based on available data, the classification criteria are not met.

- LD/LC50 values relevant for classification:		
56073-10-0 brodifacoum		
Oral	LD50	0.4 mg/kg bw (male rat and mouse)
Dermal	LD50	3.16 mg/kg bw (rat)
Inhalative	LC50/4h	3.05 mg/m³ (rat)

- **Skin corrosion/irritation** Based on available data, the classification criteria are not met.
- **Serious eye damage/irritation** Based on available data, the classification criteria are not met.
- **Respiratory or skin sensitisation** Based on available data, the classification criteria are not met.
- **Germ cell mutagenicity** Based on available data, the classification criteria are not met.
- **Carcinogenicity** Based on available data, the classification criteria are not met.
- **Reproductive toxicity** Based on available data, the classification criteria are not met.

56073-10-0 brodifacoum	
developmental toxicity	Clear developmental toxicity was not observed in rabbits or rats. However, as a precaution, Brodifacoum should be considered teratogenic to humans because it contains the same chemical moiety responsible for the teratogenicity of warfarin, a known human teratogenic agent, and it has the same mode of action that is a known mechanism of teratogenicity in humans.

- **STOT-single exposure** Based on available data, the classification criteria are not met.
- **STOT-repeated exposure** Based on available data, the classification criteria are not met.

56073-10-0 brodifacoum		
Oral	NOAEL	0.04 mg/kg bw/d (rat) The study reveals that repeated oral exposure results in toxic effects: prothrombin time prolongation, kaolin-caphalin time prolongation, haemorrhage. Based on the results of the acute dermal and inhalation toxicity studies and route-to-route extrapolation, it is justified to assume a similar concern for serious damage to health by prolonged exposure through dermal and inhalation routes also.

- **Aspiration hazard** Based on available data, the classification criteria are not met.
- **Additional toxicological information:** No further relevant information available.

- 11.2 Information on other hazards

- Endocrine disrupting properties
The mixture does not contain substances with endocrine disrupting properties in concentration equal to or greater than 0.1% by weight.

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SECTION 12: Ecological information

- 12.1 Toxicity

- Aquatic and/or terrestrial toxicity:	
56073-10-0 brodifacoum	
ErC50/72h	0.04 mg/l (selenastrum capricornutum)
EC10/6h	>0.0038 mg/l (pseudomonas putida) Based on water solubility at pH 5.2 and T=20°C.
LC50/96h	0.042 mg/l (oncorhynchus mykiss)
LC50 (diet)	0.72 mg/kg food (laughing gull)
LC50/14d	>879.6 mg/kg ww (eisenia foetida)
NOEC (reproductive toxicity)	0.0038 mg/kg food (bird)
NOEL (reproductive toxicity)	0.000385 mg/kg bw/d (bird)
NOErC	0.01 mg/l (algae)
LD50	0.31 mg/kg bw (mallard duck)
EC50/48h	0.25 mg/l (daphnia magna)

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- 12.2 Persistence and degradability

56073-10-0 brodifacoum

biodegradability	Brodifacoum is not readily biodegradable as in the test done according to the OECD 301D, 3.5% biodegradation was observed during the 28 days.
photolytic half-life	Photolytic half-life in water is 12 hours. Brodifacoum degrades rapidly by photolysis.
Hydrolitic half-life	300 days (pH 7 at 25°C). Brodifacoum is hydrolytically stable.
Persistence	The DT50 in soil is 298 days at 12°C. No data on degradation in marine water, freshwater or sediment are available. However, based on read-across with a structural analogue difenacoum, Brodifacoum is considered to be persistent and very persistent.

- 12.3 Bioaccumulative potential

56073-10-0 brodifacoum

bioconcentration factor	BCF fish = 35645 (calculated based on log Kow). BCF earthworm = 15820 (calculated based on log Kow).
bioaccumulation	For brodifacoum the screening B-criterion is fulfilled as the log Kow is above 4.5. However, BCF testing with fish would be formally required in order to be able to clarify if brodifacoum meets the B-criterion; but this test might be technically difficult to conduct as brodifacoum is highly toxic to fish. Furthermore, it should be considered that second-generation anticoagulant substances can accumulate in the liver of target rodents, and it can be assumed that they also accumulate in the livers of non-target mammals and birds. These data should be applied in addition as part of a weight of evidence approach. Based on these evidences and on information agreed on difenacoum (analogue of brodifacoum), brodifacoum should be considered bioaccumulative (B criterion fulfilled).
octanol-water partition coefficient	log Kow = 6.12 (estimated from measured Koc).

- 12.4 Mobility in soil

56073-10-0 brodifacoum

soil mobility	The substance is not considered mobile in soil.

- 12.5 Results of PBT and vPvB assessment

- PBT:

56073-10-0 brodifacoum

PBT	Brodifacoum fulfils the P, B and T criteria.

- vPvB:

56073-10-0 brodifacoum

vPvB	Brodifacoum fulfils the vP criterion.

- 12.6 Endocrine disrupting properties

The mixture does not contain substances with endocrine disrupting properties in concentration equal to or greater than 0.1% by weight.

- 12.7 Other adverse effects

56073-10-0 brodifacoum

.	The major environmental concern of Brodifacoum is primary and secondary poisoning of non-target animals.

- General notes:

Hazardous to wildlife.
Do not allow the product to reach ground water, water course or sewage system.

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SECTION 13: Disposal considerations

- **13.1 Waste treatment methods**

- **Recommendation**
At the end of the treatment, dispose uneaten bait and the packaging in accordance with local requirements.
Must not be disposed together with household garbage. Do not allow product to reach sewage system.
Do not discharge unused product on the ground, into water courses, into pipes (sink, toilets, etc.) nor down the drains.

- **Uncleaned packaging:**
- **Recommendation:** Dispose of in accordance with local requirements.

SECTION 14: Transport information

- **14.1 UN number or ID number**	
- **ADR, IMDG, IATA**	Not applicable
- **14.2 UN proper shipping name**	
- **ADR, IMDG, IATA**	Not applicable
- **14.3 Transport hazard class(es)**	
- **ADR, ADN, IMDG, IATA**	
- **Class**	Not applicable
- **14.4 Packing group**	
- **ADR, IMDG, IATA**	Not applicable
- **14.5 Environmental hazards:**	Not applicable.
- **14.6 Special precautions for user**	Not applicable.
- **14.7 Maritime transport in bulk according to IMO instruments**	Not applicable.
- **UN "Model Regulation":**	Not applicable

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SECTION 15: Regulatory information

- **15.1 Safety, health and environmental regulations/legislation specific for the substance or mixture**

- **Directive 2012/18/EU**

- **Named dangerous substances - ANNEX I** None of the ingredients is listed.

- **Seveso category** This product is not subject to Seveso directive dispositions.

- **REGULATION (EU) 2019/1021 on persistent organic pollutants (POP)**
The mixture does not contain substances identified as POP.

- **LIST OF SUBSTANCES SUBJECT TO AUTHORISATION (ANNEX XIV)**
The product does not contain any substance included in annex XIV.

- **REGULATION (EC) No 1907/2006 ANNEX XVII** Conditions of restriction: 30, 75

- **Regulation (EU) No 649/2012 (PIC)** There are no substances listed in this regulation.

- **REGULATION (EU) 2019/1148 - Explosive precursors**
The mixture does not contain explosives precursors in concentrations equal to or greater than 1%.

- **National regulations:** No further information available.

- **Other regulations, limitations and prohibitive regulations** No further information available.

- **Substances of very high concern (SVHC) according to REACH, Article 59**
The mixture does not contain SVHC substances in concentration equal to or greater than 0.1% by weight.

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- **Regulation (EU) 2024/590: substances that deplete the ozone layer**
The mixture does not contain substances that deplete the ozone layer.

- **15.2 Chemical safety assessment:**
A Chemical Safety Assessment according to Regulation (EC) No. 1907/2006 has not been carried out for the mixture.

* **SECTION 16: Other information**

This information is based on our present knowledge. However, this shall not constitute a guarantee for any specific product features and shall not establish a legally valid contractual relationship. Any responsibility derived from misuse of the product or in case of violation of current regulations is refused.

- **Relevant phrases**
EUH440 Accumulates in the environment and living organisms including in humans.
H300 Fatal if swallowed.
H310 Fatal in contact with skin.
H330 Fatal if inhaled.
H360D May damage the unborn child.
H372 Causes damage to organs through prolonged or repeated exposure.
H373 May cause damage to organs through prolonged or repeated exposure.
H400 Very toxic to aquatic life.
H410 Very toxic to aquatic life with long lasting effects.

- **Classification according to Regulation (EC) No 1272/2008**
Physico-chemical hazards: the classification of the mixture is based on the criteria established by annex I, part 2, of Regulation (EC) No. 1272/2008. If relevant, the methods are reported in section 9.
Health and environmental hazards: the classification of the mixture is based on the calculation method stated in annex I, parts 3 and 4, of Regulation (CE) No. 1272/2008, using components data.

- **Abbreviations and acronyms:**
RD50: Respiratory Decrease, 50 percent
LC0: Lethal concentration, 0 percent
NOEC: No Observed Effect Concentration
IC50: Inhibitory concentration, 50 percent
NOAEL: No Observed Adverse Effect Level
EC50: Effective concentration, 50 percent
EC10: Effective concentration, 10 percent
AEC: Acceptable Exposure Concentration
LL0: Lethal Load, 0 percent
AEL: Acceptable Exposure Limit
LL50: Lethal Load, 50 percent
EL0: Effective Load, 0 percent
EL50: Effective Load, 50 percent
ADR: Accord relatif au transport international des marchandises dangereuses par route (European Agreement Concerning the International Carriage of Dangerous Goods by Road)
IMDG: International Maritime Code for Dangerous Goods
IATA: International Air Transport Association
GHS: Globally Harmonised System of Classification and Labelling of Chemicals
EINECS: European Inventory of Existing Commercial Chemical Substances
ELINCS: European List of Notified Chemical Substances
CAS: Chemical Abstracts Service (division of the American Chemical Society)
PNEC: Predicted No-Effect Concentration (REACH)
LC50: Lethal concentration, 50 percent
LD50: Lethal dose, 50 percent
PBT: Persistent, Bioaccumulative and Toxic
SVHC: Substances of Very High Concern
vPvB: very Persistent and very Bioaccumulative
Acute Tox. 1: Acute toxicity – Category 1
Repr. 1A: Reproductive toxicity – Category 1A
STOT RE 1: Specific target organ toxicity (repeated exposure) – Category 1
Aquatic Acute 1: Hazardous to the aquatic environment - acute aquatic hazard – Category 1
Aquatic Chronic 1: Hazardous to the aquatic environment - long-term aquatic hazard – Category 1

- **References**
- Biocidal Products Committee (BPC) opinion June 2016 on the active substance;
- Assessment Report on the active substance (available at ECHA website);

- **Sources**
1. The E-Pesticide Manual 2.1 Version (2001)
2. Regulation (EC) 1907/2006 and following amendments
3. Regulation (EC) 1272/2008 and following amendments
4. Regulation (EU) 2023/707
5. Regulation (EU) 2020/878
6. Regulation (EU) 528/2012
7. Regulation (EC) 790/2009 (1st ATP CLP)
8. Regulation (EU) 286/2011 (2nd ATP CLP)
9. Regulation (EU) 618/2012 (3rd ATP CLP)

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10. Regulation (EU) 487/2013 (4th ATP CLP)
11. Regulation (EU) 944/2013 (5th ATP CLP)
12. Regulation (EU) 605/2014 (6th ATP CLP)
13. Regulation (EU) 2015/1221 (7th ATP CLP)
14. Regulation (EU) 2016/918 (8th ATP CLP)
15. Regulation (EU) 2016/1179 (9th ATP CLP)
16. Regulation (EU) 2017/776 (10th ATP CLP)
17. Regulation (EU) 2018/669 (11th ATP CLP)
18. Regulation (EU) 2019/521 (12th ATP CLP)
19. Regulation (EU) 2018/1480 (13th ATP CLP)
20. Regulation (EU) 2020/217 (14th ATP CLP)
21. Regulation (EU) 2020/1182 (15th ATP CLP)
22. Regulation (EU) 2021/643 (16th ATP CLP)
23. Regulation (EU) 2021/849 (17th ATP CLP)
24. Regulation (EU) 2022/692 (18th ATP CLP)
25. Regulation (EU) 2023/1434 (19th ATP CLP)
26. Regulation (EU) 2023/1435 (20th ATP CLP)
27. Regulation (EU) 2024/197 (21st ATP CLP)
28. Regulation (EU) 2024/2564 (22nd ATP CLP)
29. Directive 2012/18/EU (Seveso III)
30. ECHA web site

- *** Data compared to the previous version altered.**