Procedure file

Basic information

RSP - Resolutions on topical subjects	2020/2852(RSP)	Procedure completed
Resolution on the draft Commission implementing regulation approving carbendazim as an existing active substance for use in biocidal products of product-types 7 and 10 Subject 3.40.01 Chemical industry, fertilizers, plastics 3.70.13 Dangerous substances, toxic and radioactive wastes (storage, transport) 4.60.08 Safety of products and services, product liability		

Key players

European Parliament	Committee responsible	Rapporteur	Appointed
	ENVI Environment, Public Health and Food Safety		27/10/2020
			27/10/2020
		S&D ARENA Maria	27/10/2020
			27/10/2020
		RIVASI Michèle	
		HAZEKAMP Anja	
		NI EVI Eleonora	

Key events

26/11/2020	Results of vote in Parliament		
26/11/2020	Decision by Parliament	T9-0326/2020	Summary

Technical information

Procedure reference	2020/2852(RSP)
Procedure type	RSP - Resolutions on topical subjects
Procedure subtype	Resolution on implementing act or powers
Stage reached in procedure	Procedure completed
Committee dossier	ENVI/9/04498

Documentation gateway

Motion for a resolution		B9-0366/2020	24/11/2020	EP	
Text adopted by Parliament, single reading		T9-0326/2020	26/11/2020	EP	Summary

Commission response to text adopted in plenary		SP(2021)129	02/06/2021	EC	

Resolution on the draft Commission implementing regulation approving carbendazim as an existing active substance for use in biocidal products of product-types 7 and 10

The European Parliament adopted by 458 votes to 219, with 19 abstentions, a resolution objecting to the draft Commission implementing regulation approving carbendazim as an existing active substance for use in biocidal products of product-types 7 and 10.

The draft Commission implementing regulation seeks to approve carbendazim as an existing active substance for use in biocidal products of product-type 7 (film preservatives) and product-type 10 (masonry preservatives), for a period of three years.

However, Parliament considered that the draft Commission implementing regulation is not consistent with EU law, in that it is not compatible with the aim and content of Directive 98/8/EC or Regulation (EU) No 528/2012.

Furthermore, it considered, in view of:

- the hazardous properties of carbendazim,

- its environmental fate, as well as the lack of risk management measures stated in the supporting documents,

- the lack of data to decisively conclude on the absence of suitable alternatives,

- the seven-year period that has passed since the submission of the assessment reports, and

- the lack of coherence between the Commission decisions on the uses of carbendazim in product-types 7, 9 and 10,

that the draft Commission implementing regulation to approve carbendazim as an existing active substance for use in biocidal products of product-types 7 and 10, even for a short period of three years, is not proportionate in light of the unacceptable risks it poses to human health and the environment, and should have lead the Commission to the conclusion of unacceptable risks, as the use of carbendazim in a product still gives rise to concerns.

It called on the Commission to withdraw its draft implementing regulation and to submit a new draft to the committee, proposing not to approve carbendazim as an active substance for use in biocidal products of product-types 7 and 10.