Basic information	
2020/2852(RSP) RSP - Resolutions on topical subjects	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 events			
Date	Event	Reference	Summary
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

European Parliament

Document type	Committee	Reference	Date	Summary
Motion for a resolution		B9-0366/2020	24/11/2020	
Text adopted by Parliament, single reading		T9-0326/2020	26/11/2020	Summary

European Commission

Document type	Reference	Date	Summary
Commission response to text adopted in plenary	SP(2021)129	02/06/2021	

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

2020/2852(RSP) - 26/11/2020 - Text adopted by Parliament, single reading

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.