Extension of Authorisation Number: 1571 of 2024

EXTENSION OF AUTHORISATION FOR A MINOR USE OF A PLANT PROTECTION PRODUCT

PLANT PROTECTION PRODUCTS REGULATION No 1107/2009

Extent of authorisation:	Great Britain and Northern Ireland
Product name*:*	Kenja
Active ingredient*:*	400 g / l Isofetamid
MAPP number*:*	20200
Product authorisation holder*:*	ISK Biosciences Europe NV (Registered Company no. 601473)
Marketing company*:*	Certis Belchim B.V.

This Extension of authorisation ends: on the final expiry date of use for the authorised product

If the authorisation of the above product is withdrawn or amended, this Extension of authorisation will end on the same date as the authorisation for the product.

This Extension of authorisation will be withdrawn or amended before its end date if any of the active substances contained in the product are withdrawn from the GB Approvals Register or list of approved active substances included in Regulation (EU) No 540/2011, or if a decision is taken to withdraw or amend this Extension of authorisation under Regulation No 1107/2009 on any other grounds.

The circumstances in which this Extension of authorisation will be withdrawn or amended are set out in Regulation No 1107/2009.

This extension of authorisation for minor uses applies to all authorised parallel trade products issued under Article 52 of Regulation No 1107/2009 for which Kenja with MAPP 20200 is the reference product.

Digitally signed by: Stephen Thornley
Date: 2024.10.16 11:34:45 Z

HSE Digital Signature

This and the attached Appendices 1 and 2 are signed by the Health and Safety Executive for and on behalf of the Secretary of State, the Welsh Ministers, the Scottish Ministers and the Department of Agriculture, Environment and Rural Affairs in Northern Ireland.

Date of issue: 16 October 2024

EXPLANATORY NOTES

1. This is Extension of authorisation number 1571 of 2024.

2. This Extension of authorisation will be published on HSE's website.

3. Application reference number: COP 2022/00990

4. Persons using the product to which this Extension of authorisation applies should acquaint themselves with and observe all requirements contained in the Regulation No 1107/2009, including the duty on the holder of any Extension of authorisation to notify information on potentially dangerous effects, a contravention of which is a criminal offence under those Regulations.

5. Neither the efficacy nor the phytotoxicity of the product for which this Extension of authorisation has been granted has been assessed and, as such, the user bears the risk in respect of failures concerning its efficacy and phytotoxicity.

6. In this notice Regulation No 1107/2009 means:
   In relation to Great Britain, Assimilated Regulation No 1107/2009 as it has effect in Great Britain.
   In relation to Northern Ireland, Regulation (EC) No 1107/2009 as it has effect by virtue of the Protocol on Ireland/Northern Ireland in the EU withdrawal agreement.

7. In this notice Regulation (EU) No 540/2011 means:
   In relation to Northern Ireland, Regulation (EU) No 540/2011 as it has effect by virtue of the Protocol on Ireland/Northern Ireland in the EU withdrawal agreement.

ADVISORY INFORMATION

IMPORTANT: When applying this product under the terms of this Extension of authorisation, comply with any resistance guidance or restrictions stated on the product label.
Total reliance on one pesticide will hasten the development of resistance. Pesticides of different chemical types or alternative control measures should be included in the planned programme. Alternating with different modes of action is a recognised anti-resistance strategy.

This Extension of authorisation relates to the use of ‘Kenja’ (M20200) for the control of botrytis (*Botrytis cinerea*) sclerotinia (*Sclerotinia sclerotium*) and *Monilinia*, to be applied by via conventional horizontal boom sprayer, handheld tank and lance sprayer or handheld knapsack sprayer in 200 to 500 litres water per hectare, all as a downward spray. It is for use on protected (both permanent/glasshouse and temporary/polytunnels) container-grown ornamental plant production (pot and bedding, hardy nursey stock, fruit nursery trees, Christmas trees) once per crop based on a maximum of three crop cycles per year in any single protected structure. And on protected (both permanent/glasshouse and temporary/polytunnels) soil-grown ornamental plant production (non-bulb cut flowers, bulb cut flowers (including rhizomes, corms and tubers) once per year.

Botrytis cinerea strains resistant to SDHI fungicides are known to occur. Where such strains occur, the level of ‘Kenja’ efficacy may be adversely affected.’
Refer to FRAC for guidance on the use of SDHI fungicides in mixtures and for information on the maximum number of SDHI applications permitted on each crop per season

APPENDIX 1: CONDITIONS OF EXTENSION OF AUTHORISATION

The conditions below are obligatory. They must be complied with when the Extension of authorisation occurs. Failure to comply with the following conditions will be likely to result in the withdrawal or amendment of the Extension of authorisation under Regulation No 1107/2009 and may result in other enforcement action, including prosecution. For the purposes of this Extension of authorisation only, the conditions and/or requirements shown below supersede any corresponding conditions and/or requirements set out on the label or otherwise provided for under the product authorisation **which would otherwise apply**.

Use:

Field of use: **ONLY AS A FUNGICIDE**

User: Professional

Crops/situations:	Maximum individual dose: (litres product / ha)	Maximum total dose:	Maximum number of treatments: (per year)	Latest time of application:
Protected ornamental plant production - container grown crops	1	-	1 (See Other Specific Restriction 3)	BBCH 87
Protected ornamental plant production	1	-	1 (See Other Specific Restriction 4)	BBCH 87

Operator Protection:

(1) Vehicle-mounted or trailed horizontal or vertical boom sprayers must only be used where the operator's normal working position is within a closed cab with a suitable in-cab filtration system* or suitable respiratory protective equipment** must be worn during application in protected situations. *Closed cabin meeting at least EN 15695 category 3. **Disposable filtering facepiece respirator to at least EN149 FFP3 or equivalent.

(2) Other engineering control of operator exposure must also be used where reasonably practicable in addition to the following personal protective equipment:

(a) Operators must wear suitable protective clothing (coveralls) and suitable protective gloves when handling contaminated surfaces or contact with treated foliage is unavoidable during application in protected situations.

(b) Operators must wear suitable protective gloves when handling the concentrate.

(c) Workers must wear suitable protective clothing (in which arms, body and legs are fully covered) and suitable protective gloves* when handling protected treated crops, re-entering protected treated areas within 11 weeks after treatment. *Meeting at least glove safety standard EN374-2:2019, Level 2 and CE category III. Such gloves can be identified by a CE Mark with four digits below, and the EN374 pictogram for micro-biological hazards.

(3) However, other engineering controls in addition to those specified above may replace personal protective equipment if a COSHH assessment shows they provide an equal or higher standard of protection.

Environmental protection:

When used under permanent protection on crops grown in synthetic media please note the following:

The use of this product in recirculating water systems in a glasshouse may result in dilute pesticide waste that requires disposal. All dilute pesticide waste must be disposed of safely and legally to protect humans, wildlife and the environment, especially groundwater and surface water. Pesticide disposal advice is detailed in the ‘Code of Practice for Using Plant Protection Products (Section 5: Disposing of Pesticide Waste).

Other specific restrictions:

(1) This product must only be applied in accordance with the terms of this extension of authorisation, the product label and/or leaflet and any additional guidance on extensions of authorisation.

(2) Managers must carry out a thermal comfort checklist (see - http://www.hse.gov.uk/temperature/assets/docs/thermal-comfort-checklist.pdf) prior to worker re-entry tasks. If needed, an additional heat stress check list

and associated risk assessment must be undertaken (see-
http://www.hse.gov.uk/temperature/assets/docs/heat-stress-checklist.pdf) and the records retained. Temperature and humidity inside tunnels/greenhouses should be monitored during re-entry tasks. If conditions become such that there is a risk of heat related illness, or workers complain of ill effects, then work must cease until the risk is reduced. It is not acceptable for workers to remove clothing and continue working.

(3) When applied to container grown ornamentals this product must only be applied three times a year in any protected structure (once per crop on a maximum of three crop cycles per year).

(4) For use only on non-bulb cut flowers, bulb cut flowers (including rhizomes, corms and tubers), forest nursery, Christmas tree production and fruit nursery trees.

APPENDIX 2: GENERAL CONDITIONS FOR AN EXTENSION OF AUTHORISATION

Failure to comply with the following conditions will be likely to result in the withdrawal or amendment of the Extension of authorisation under Regulation No 1107/2009 and may result in other enforcement action, including prosecution.

Adverse effects:

The authorisation holder must immediately notify the Secretary of State, the Welsh Ministers, the Scottish Ministers and the Department of Agriculture, Environment and Rural Affairs in Northern Ireland (care of the Health and Safety Executive), if they have any new information on the potentially adverse effects of the authorised product, or of residues of an active substance in that product when used in accordance with the conditions of this authorisation. For those products authorised under Regulation (EC) No 1107/2009 as it has effect by virtue of the Protocol on Ireland/Northern Ireland in the EU withdrawal agreement, authorisation holders must also tell the other relevant competent authorities of the EC Member States (a list of which is available from the Health and Safety Executive) and the EC Commission. Failure to comply with this requirement is an offence.

Provision of information:

The authorisation holder must comply with all requests for information required by, or on behalf of, the Secretary of State, the Welsh Ministers, the Scottish Ministers or the Department of Agriculture, Environment and Rural Affairs in Northern Ireland in accordance with Regulation No 1107/2009.