# Azadirachtin

## Status under Reg. (EC) No 1107/2009 (repealing Directive 91/414/EEC)

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| **Status:** | Approved | **Current Legislation:** | 2011/44/EU , Reg. (EU) No 540/2011 |
| **Date of approval:** | 01/06/2011 | **Expiration of approval:** | 31/05/2021 |
| **RMS:** | DE | **EFSA Risk Assessment:** | ✓ |
| **Category:** | IN | **Review Report:** | |

**Remarks:** Initially non included by Decision 2008/941. Included as from 1 June 2011 following re-submission for inclusion according to Reg. 33/2008.

## Classification

**No classification**

## Authorisations at national level

| | | | |
|---|---|---|---|
| **Authorised in:** | AT, BE, BG, CY, CZ, DE, EE, ES, HU, IT, LT, LU, LV, NL, PT, SI | **In progress for:** | SE |

## Toxicological information

Azadirachtin

| ADI: | Source: | Remark: | ARfD: | Source: | Remark: | AOEL: | Source: | Remark: |
|---|---|---|---|---|---|---|---|---|
| 0,1 | 11/44/EU | | 0,75 | 11/44/EU | | 0,1 | 11/44/EU | |
| **Other:** | | | | | | | | |

Where no units are shown, values are expressed in mg/kg bw/day

## EU - Maximum Residue Levels (Reg. (EC) No 396/2005) (MRLs)

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|---|---|---|---|
| **Legislation:** | **Azadirachtin**<br>Reg. (EC) No 149/2008 | **Annexes:** | **Azadirachtin**<br>Annex IIIA |

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