

Recommendations

In this guide

[In this guide](#)

1. [Introduction and Background - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
2. [Summary of 2025 EFSA draft evaluation - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
3. [Acute toxicity studies - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
4. [Repeat dose toxicity studies - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
5. [Observations in Humans - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
6. [Mode of action - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
7. [Occurrence data and dietary exposure assessment for the European population - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
8. [Risk characterisation - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
9. [Uncertainty analysis - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
10. [Recommendations - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
11. [Questions on which the views of the Committee are sought - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
12. [List of Abbreviations - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)
13. [References - EFSA draft scientific opinion on risks for human health of plant lectins in food](#)

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35. EFSA recommended that for the quantification of active and non-active lectins, the development and validation of analytical methods was needed. In addition to this, studies using PHA were needed for information on absorption, distribution, metabolism, and excretion (ADME), immunotoxicity, and gastrointestinal endpoints in humans and rodents. Further information was also required for an in-depth exposure assessment.