

Commission

Approval of pesticides and herbicides in the EU: *How does it work* ?

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The approval of active substances (including for herbicides) in the European Union is a rigorous process aimed at ensuring the protection of plants against pests and diseases, as well as the safety of human health and the environment. The process involves a rigorous scientific assessment by experts from the Member States and European agencies, which forms the basis for a decision on whether the substance can be used in plant protection products. The **comitology procedure** is a key element in the final step of the decision-making process

APPLICATION SUBMISSION:

Companies who want to market pesticides containing a specific active substance need to submit an application for approval to a Member State (or a group of Member States), which then acts as Rapporteur Member State(s).

RAPPORTEUR MEMBER STATE(S) ASSESSMENT:

The Rapporteur Member State (or States) conducts a comprehensive evaluation of the dossier for the active substance.

The evaluation includes a risk assessment for human health and the environment, including, residues in food.

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COMPANY X

PEER REVIEW PROCESS:

The assessment is peer-reviewed by experts from all other Member States and EFSA (European Food Safety Authority) to ensure it is scientifically robust and all data are considered.

EFSA finalises its opinion (in the form of a Conclusion), based on the peer review. This is then transmitted to the European Commission, which prepares a draft review report and a draft Regulation on the approval of the active substance.

STANDING COMMITTEE DELIBERATION:

The draft Regulation for approval or non-approval is prepared by the Commission and discussed in the Standing Committee on Plant Protection Products.

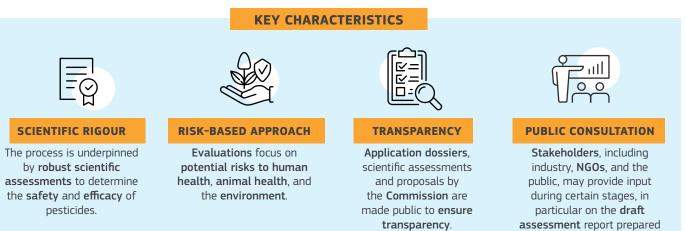
The Standing Committee is composed of representatives of all EU countries and votes on the Commission proposal.





After the Committee delivers a positive opinion, or if the Appeal Committee delivers no opinion, the decision is adopted by the Commission. **A new active substance is usually approved for a maximum of 10 years** (if it is "low risk" it is approved for 15 years), while a renewed approval can be granted for up to 15 years. Approval with restrictions is also possible.

After the approval of an active substance, companies wishing to place products containing the active substance on the market must **seek authorisation in the Member States**, who must again assess the safety of each plant protection product for each use prior to granting an authorisation.





https://commission.europa.eu/law/law-making-process/adopting-eu-law/implementing-and-delegated-acts/ comitology_en

by the Rapporteur Member State.

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