



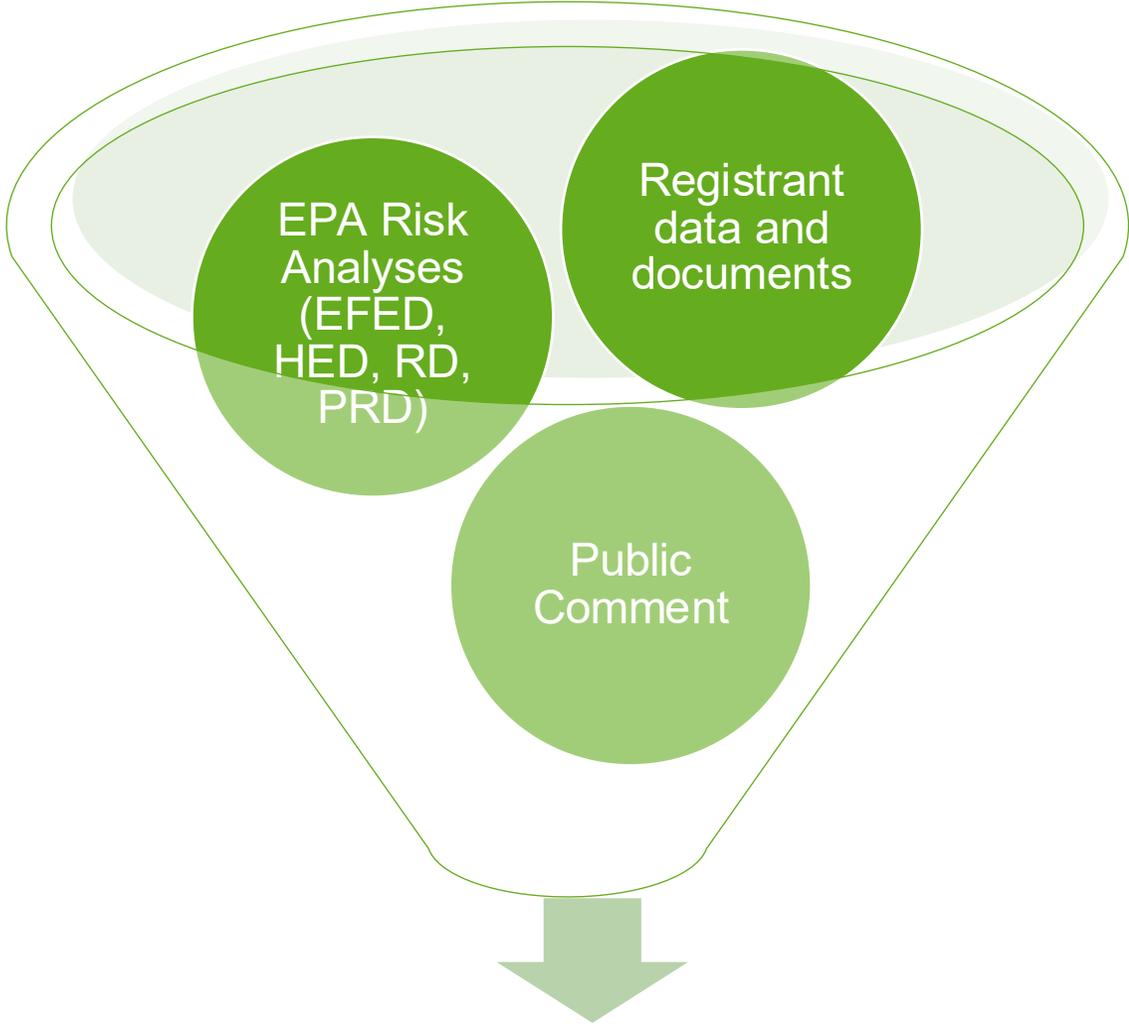
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Registrations: Registrant Perspective

March 2, 2026



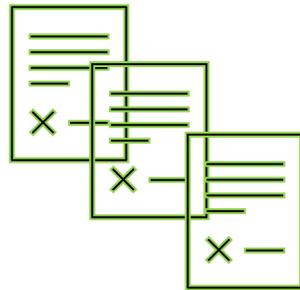
Overview of Process



**EPA Registration of
Pesticide Product**

Registrant Submissions

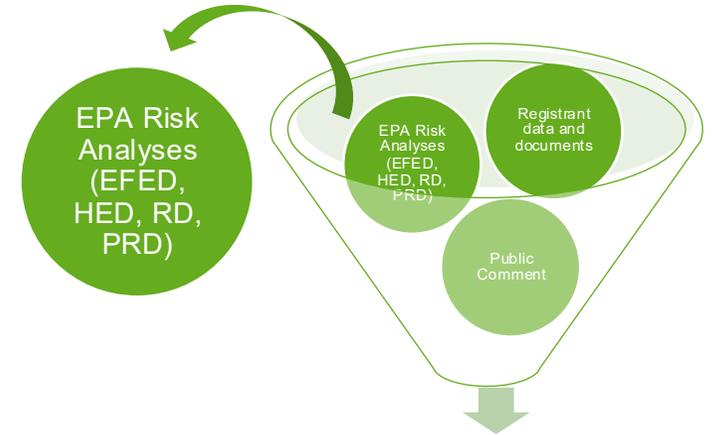
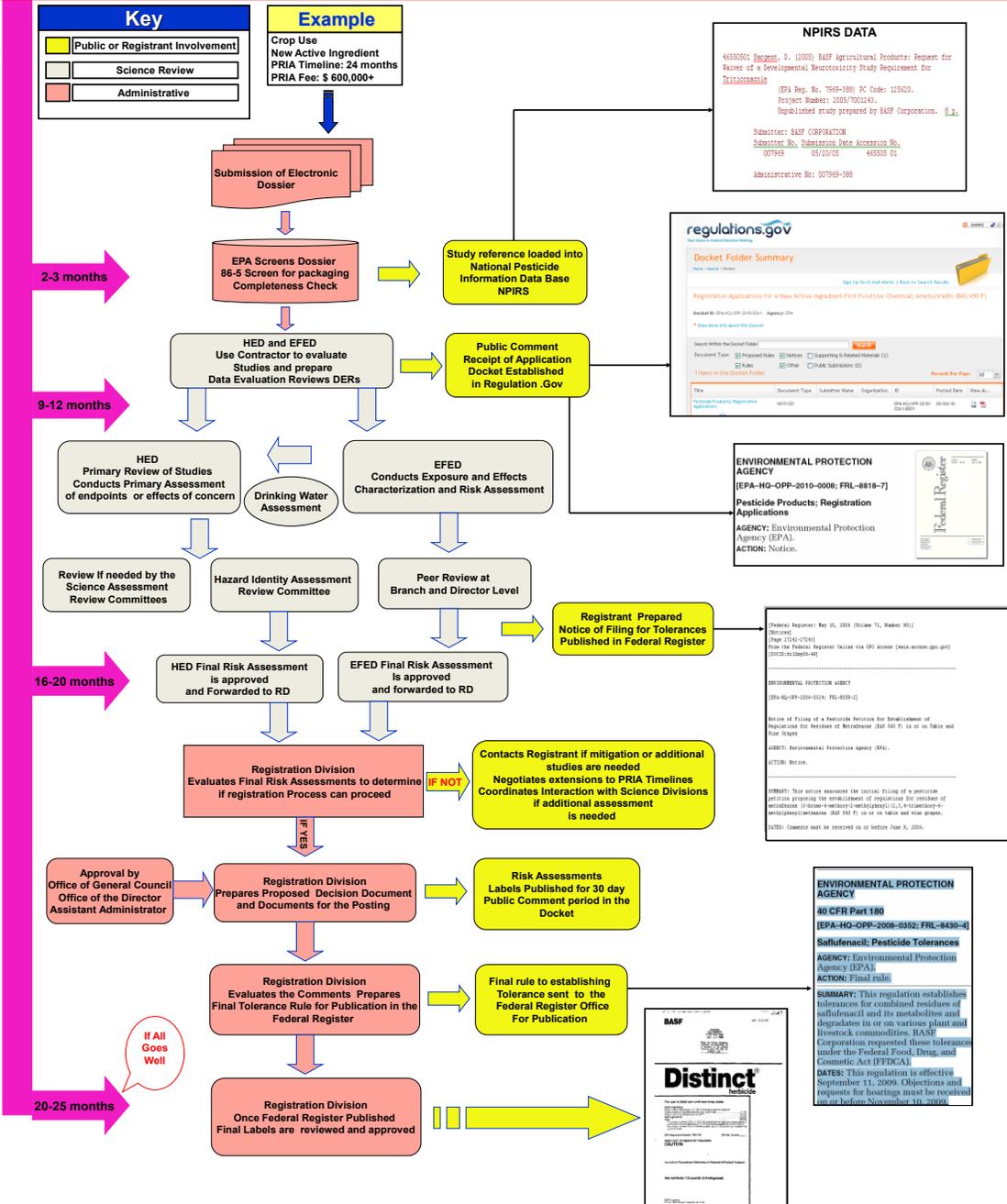
- >200 studies, including
 - ▶ Environmental fate
 - ▶ Ecotoxicological data
 - ▶ Impact on nontarget species
 - ▶ Residue chemistry
 - ▶ Human health toxicity data
- Benefits document
- Proposed label
- PRIA fee when applicable



EPA Risk and



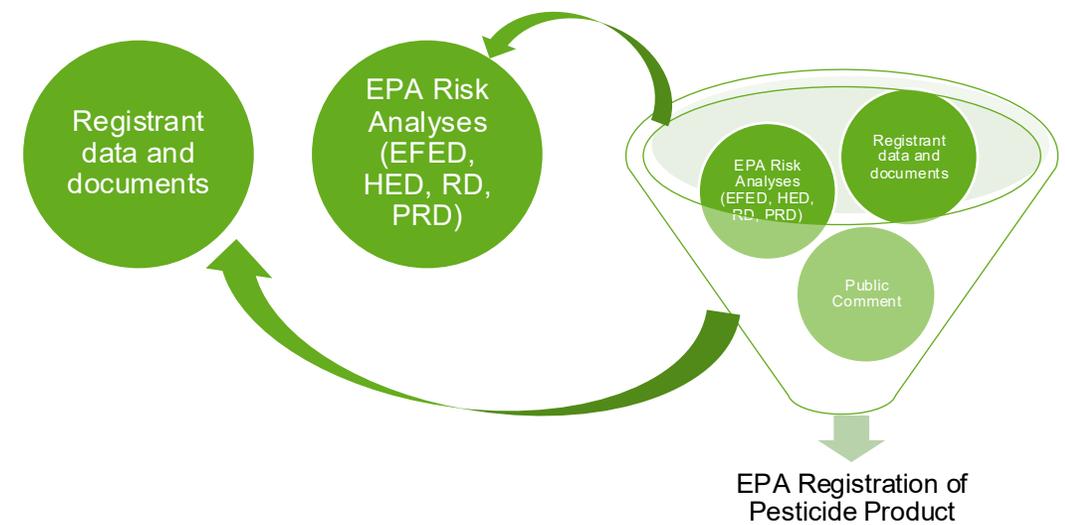
US EPA Regulatory Process



EPA Registration of Pesticide Product

Label Development

- Registrants provide the draft label
 - ▶ EPA Label Review Manual
 - ▶ Weed resistance management language (herbicides)
 - ▶ Use Directions and expertise on the end use product
- EPA RD/PRD manager and the registrant will confer throughout the process
 - ▶ During the assessments – clarification, data questions to the registrant
 - ▶ After the assessment – label language clarification



Public Input

Certain actions will require public comment:

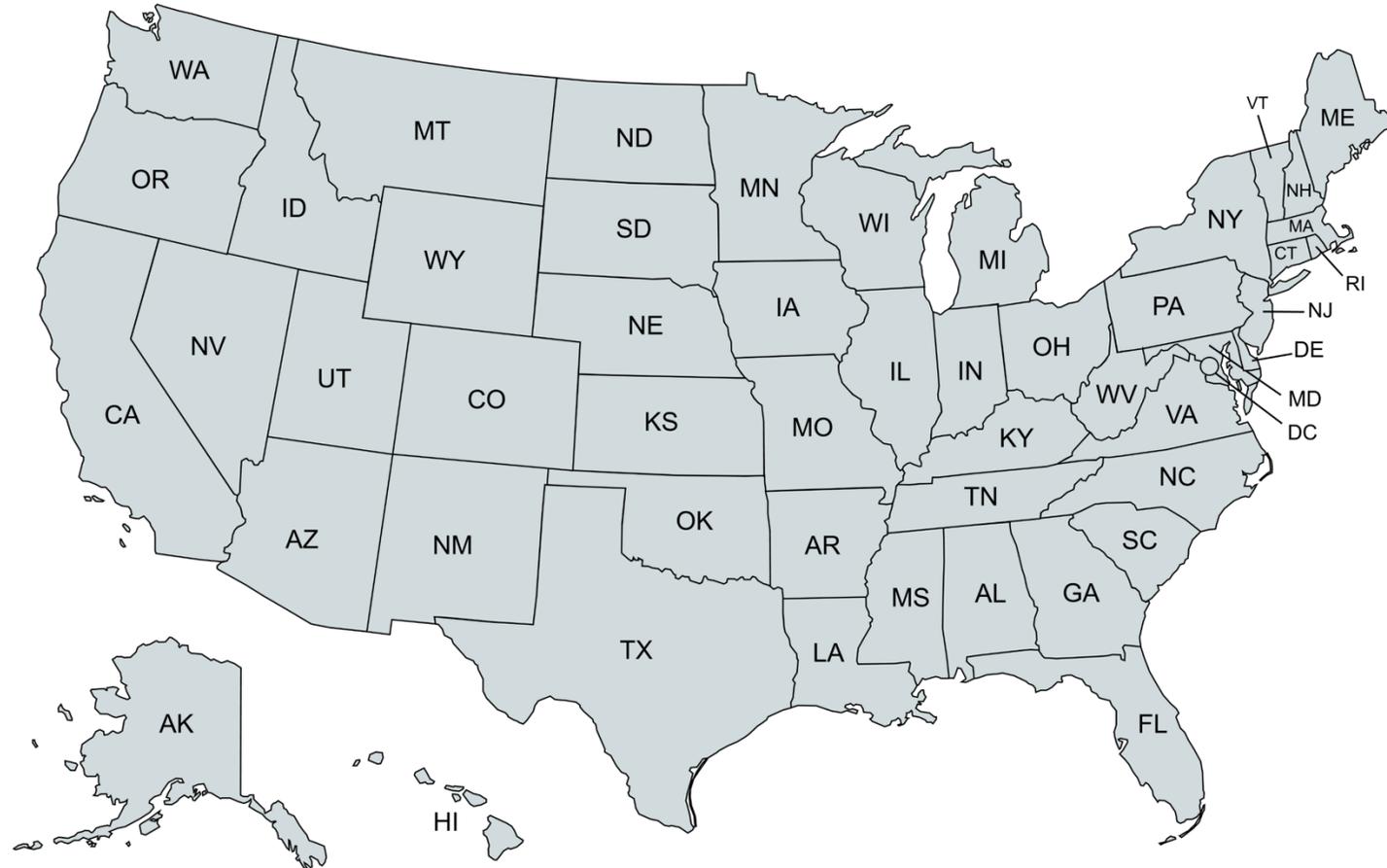
- ▶ New registrations
- ▶ New uses
- ▶ Tolerances
- ▶ Cancellations
- ▶ Some risk assessments
- ▶ Proposed interim/final decisions



State Submissions

Submission strategy is up to the registrant:

- ▶ Where:
 - Dependent on market for product
 - Nationwide approach
- ▶ When:
 - EPA 18 month rule
 - Product launch plans
 - Timing of next production of product
- ▶ How:
 - Paper
 - State specific portal
 - STAR
 - Electronic
 - Combination



Areas for Improvement

- PRIA timelines – predictability and certainty for all parties
- Structured labeling – consistency
- PR 98-10 – draft on notifications and non-notifications available for comment



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