

The New Pest Control Products Act



- Parliamentary Process
- Objectives of the Amendments
- Considerations in Drafting Bill
- Main Elements of New PCPA
- Benefits

Parliamentary Process

- First Reading
- Second Reading
- Committee study and report
- Third Reading
- Senate
- Royal Assent
- *Regulations

Objectives of the Amendments

- * Health and environmental protection
- Openness and transparency
- Cost-effective regulation
- PRR recommendations
- Good public policy

Considerations in Drafting Bill

- Appropriate role of federal government
 - national registration system
 - prevent unacceptable risks
 - avoid unsupported claims
 - facilitate sustainable development

Considerations in Drafting Bill (cont'd)

- Importance of pesticides in pest management and the economy
- Need to minimize risks and encourage sustainable pest management
- Cost-effective and open regulation

Considerations in Drafting Bill (cont'd)

- Divergent interests and right to participate
- Cooperation among federal departments
- Partnership relationship with provinces and territories
- International obligations

Risk Management Approach

* Risk

products posing unacceptable risk not registered

Value

- efficacy
- sustainable pest management
- competitiveness
- Transparency / public participation

Mandatory Reporting of Adverse Effects

Statute

- basic obligation to report
- basic process for handling information
- consequences
- Details in regulations
 - > who
 - > what
 - > when

Re-evaluation and Special Review

- * Re-evaluation
 - triggered by change in information requirements or evaluation procedures
- Special Review
 - triggered by specific concerns
 - public may request
- Consequences
 - confirmation, amendment or cancellation

Export Restrictions

Statute

export of products of concern by permit only

Regulations

- criteria for designation as "product of concern"
- requirements to obtain permit

Enforcement

- Clarify prohibitions
- Increase penalties
- * Modernize powers of inspectors
- Ensure authority to prescribe circumstances and conditions to allow:
 - tank mixing
 - lower than label rates



Public Consultation

- Major registration decisions
 - new active ingredients
 - major new uses
 - re-evaluations and special reviews
- Proposed Regulatory Decision Documents (PRDDs)
 - summaries of risk and value assessments

Access to Test Data

- Provide access to data and evaluations supporting pesticide registrations
- Prevent:
 - "unfair commercial use" of data
 - disclosure of confidential business information

Access to Test Data (cont'd)

- Ensure that confidential information can be shared when appropriate
 - e.g., with provincial/territorial regulators
- Public registry to facilitate access to non-confidential information
- * Details in regulations to allow:
 - consultation during development of regulations
 - adaptation in future as policies evolve

Public Access to Review Mechanisms

- Public may request:
 - special review of registered pesticides
 - reconsideration of registration decisions
- Transparent review processes
- Can lead to confirmation, amendment or cancellation

Regulation-making Authorities

- Registration types
- Research permits
- Lower than label rates
- National pesticide database
- Good laboratory practices
- Data protection





Public, Provinces/Territories, Stakeholders

- safe pesticides
- more opportunities to influence registration decisions
- access to information supporting registration decisions
- increased public confidence in federal registration decisions

Benefits (cont'd)

Provinces / Territories

- easier access to safety information
- flexible regulations
 - e.g., lower than label rates
- more effective collaboration in compliance and enforcement

Benefits (cont'd)

Manufacturers

- international harmonization leads to lower costs and timely registration decisions
- protection from unfair commercial use of data

Benefits (cont'd)

Users

- international harmonization leads to timely access to sustainable pest management tools and level-playing field
- increased confidence in agri-food and other products



- Legislative proposals based on PRR recommendations
- Awaiting direction from Minister of Health
- Amendments benefit all stakeholders
- Details in regulations will be developed in consultation with stakeholders