How APHIS, EPA, and FDA may Use Study Findings

Findings of report will be helpful to inform ongoing and future agency activities, incl. development of long-term strategy (Ref: 2015 EOP memo). USDA/APHIS, EPA, and FDA may use the report to:

- Gain a better understanding of future products and how they fit within U.S. regulatory system
- Consider any necessary updates to scientific assessments
- Consider any necessary updates to regulatory processes or procedures
- Help enhance communication with stakeholders

APHIS, EPA, and FDA may Use Study Findings to . . .

- Gain a better understanding of future products (incl. those potentially developed in other countries), and consider how products fit within:
  - individual agency regulatory frameworks; and
  - broader U.S. Coordinated Framework

- Consider any necessary updates to scientific assessments
  - Any refinements to current risk/safety assessment frameworks, considering risk to humans, animals, plants, or environment due to new or different:
    - Properties
    - Routes of exposure
    - Hazards
    - Other risk-related characteristics
  - Plan regulatory science activities

USDA

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

FDA U.S Food and Drug Administration

APHIS, EPA, and FDA may Use Study Findings to . . .

- Consider any necessary updates to regulatory processes or procedures
  - Any new or revised safety testing methods or data needed in regulatory submissions
  - Any need for new/updated recommendations or guidance for industry
  - Any additional expertise (hiring) needed for regulatory review of future products

- Help enhance communication with stakeholders
  - Identify any additional relevant stakeholders
  - Plan public engagement and outreach activities