



# What is the advantage of dismantling FDA and EPA on human health

Reports indicate that the dismantling or reduction of the Food and Drug Administration (FDA) and Environmental Protection Agency (EPA) could potentially be driven by a desire to reduce perceived bureaucratic burdens and lower costs for businesses. [↗](#)

However, the consequences of such actions are widely reported to be detrimental to human health and the environment:

- **Weakened Safeguards and Increased Pollution:** Weakening or eliminating regulations imposed by these agencies is projected to lead to increased pollution of air, water, and land, potentially impacting vulnerable communities disproportionately. This could result in higher rates of pollution-related illnesses and premature deaths.
- **Reduced Oversight of Products and Increased Risk:** The FDA is responsible for ensuring the safety and effectiveness of food, drugs, and medical devices. Reductions in staff and funding could lead to decreased inspections, slower approvals of new drugs, and potentially an increase in unsafe products reaching consumers.
- **Setbacks in Environmental Protection and Public Health Programs:** Dismantling or reducing these agencies could reverse progress made in areas like protecting communities from lead poisoning and addressing climate change.
- **Negative Economic Impacts:** While some proponents argue for cost savings, studies suggest that the economic benefits of environmental regulations often outweigh the costs by preventing illnesses, premature deaths, and supporting industries like clean energy. [↗](#)

In essence, while the argument for dismantling these agencies might center on deregulation and cost reduction for businesses, many sources suggest that the potential benefits to human health are significantly outweighed by the risks associated with diminished oversight and environmental protection. [↗](#)

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↪ What historical events led to the creation of the FDA and EPA?

↪ Provide examples of potential impacts of environmental deregulation on specific industries

↪ How could FDA staff cuts affect drug approvals and safety?

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