

# A New Canadian Federal Approach to Health

## A Provincial Opportunity and a National Necessity

The implications of an aging population on health systems and succession of Canadian business create two key challenges for Canada's provinces and territories:

- Healthcare costs (including addiction crises, product shortages and delays in access to medicines) and
- The cost of living/doing business (including increasing demand for Canadian made goods to address insecurities in Canada's supply chain).

The challenges above are directly related to the federal approach to health including:

- Establishing the legal (criminal) benchmark for making and selling products in Canada
  - Canada's unique regulatory approaches that do not align internationally and are not validated against health outcomes.
- Establishing federal health research and investment priorities and the related administration
  - Canada's disparities in the priorities of health charities the federal departments who jointly fund health research as well as the laws that limit their respective overhead.
- Ensuring that Canadian compliant business practices are competitive and easy to start up/grow
  - Canada's ad-hoc and isolated efforts to validate federal laws against enforcement resources or maintain consistency in the application of laws across retail settings online and in-person.

### What is the federal approach to health?

Essentially, a group of departments and agencies reporting to the Minister of Health who, together, set national health program priorities, and most importantly, establish national standards using criminal law power. Standards, that govern access to healthcare products, research activities, farming practices, and the manufacturing of every personal commodity sector ranging from household appliances and furniture to food, medicine, pesticides and recreational substances.

The use of criminal law power has direct implications on the priorities and coordination of policing forces at all levels of government. However, this power largely does not extend what citizens can make themselves or import for personal use. As it relates to access to medicines as well as the costs of business and everyday items, the government has increasingly struggled with combining risk observations and good intentions into to presumptive-risk-based regulatory controls. Effectively, introducing paternalistic approaches into criminal-law-based public protection.

Criminal law power was never designed for the management of presumptive risks - those that cannot be validated nor measured. Fair competition, investment, efficient operational models, enforcement priority setting and innovation are all hindered by federal approaches that have been based on overprotective regulations.

It is critical to recognize that disparities between the regulatory approaches over the sale and manufacture of something, versus something that can be made at home, can erode public trust by creating a double standard of risk-tolerance between the people who buy things versus those who make things.

The growth of this double standard can likewise incubate a form of *divisiveness in the real world between people who buy things and those who make things.*

**Canada's federal approach to health is not calibrated to a world of shifting and growing risk tolerances.**

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