Eurofins BioPharma – Regulatory Services

Eurofins BioPharma Toronto provides a wide array of professional regulatory services. With years of industry-leading experience, we can take your company from concept to operation in compliance with Health Canada for products such as drugs, medical devices, natural health products, and cosmetics.

Pharmaceuticals for sale in Canada must be properly labelled, licenced, manufactured and stored in compliance with all legislative requirements. We help our clients establish a well-designed and correctly implemented pharmaceutical quality system (ie. Quality Management System) that incorporates good manufacturing practices (GMP) and quality risk management.

Services provided by our experienced consultants include:

- Regulatory classification of products based on ingredients and label claims
- Product or Establishment Licence Applications,
- Amendments and Notifications
  - Pharmaceutical Drug Products (DIN)
  - Natural Health Products (NPN, DIN-HM)
  - Medical Devices (MDL)
- Product label development to help maximize claims and expedite product approvals
- Cosmetic product consulting (label & ingredient reviews, cosmetic notification)
- Strategic planning, design of a GMP compliant facility and
- Development and implementation of a quality system (GMP)
- Good Pharmacovigilance Practices (GVP) services

Shall we guide you through the regulatory pitfalls?

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