MDCCD 2022-2025 STRATEGIC PLAN

WHO WE ARE

MDCCD MANAGES AND DELIVERS NATIONAL COMPLIANCE AND ENFORCEMENT PROGRAMS FOR MEDICAL DEVICES; BLOOD; DONOR SPERM AND OVA; CELLS, TISSUES AND ORGANS; CLINICAL TRIAL AS WELL AS MONITORING AND ENFORCING THE IMPORTATION OF HEALTH PRODUCTS AT THE BORDER.

VISION

TO BE A WORLD-CLASS COMPLIANCE AND ENFORCEMENT ORGANIZATION WITH LEADING EDGE PRACTICES WHICH RESULTS IN VISIBLE, MEASURABLE AND POSITIVE RESULTS TO PROTECT CANADIANS, THROUGH AN ENGAGED AND HEALTH WORKFORCE

MISSION

TO BE A COMPLIANCE AND ENFORCEMENT LEADER THAT INFORMS AND PROTECTS CANADIANS FROM HEALTH RISKS ASSOCIATED WITH VARIOUS HEALTH PRODUCTS

STRATEGIC OBJECTIVES



TRANSFORMATION

Transform and improve our C&E approach to become more agile, consistent and risk based



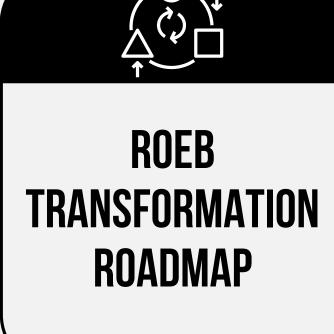
MODERNIZATION

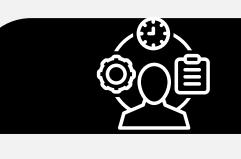
Establish an efficient and responsive approach to program delivery

PEOPLE AND WORKPLACE

Create and support a respectful, resilient and thriving workplace

GUIDING PRINCIPLES





CHANGE MANAGEMENT PRINCIPLES



ROEB WELLNESS CHARTER

PILLARS

TRANSFORMATION

TRANSFORM AND IMPROVE
OUR C&E APPROACH TO
BECOME MORE AGILE,
CONSISTENT AND RISK BASED

OBJECTIVES

Objective 1: Maximise operational efficiency through the strategic use of resources, partnerships, and process automation and optimization

Objective 2: Implement data analytics, enhanced risk-based activities and intelligence gathering to inform operational and strategic decisions

Objective 3: Enhance program and regulatory agility to better respond to emerging global and domestic issues

MODERNIZATION

ESTABLISH AN EFFICIENT AND RESPONSIVE APPROACH TO PROGRAM DELIVERY

Objective 1: Centralise key horizontal activities to better support programs

Objective 2: Improve operational processes to be more effective and efficient

Objective 3: Improve IT systems to facilitate program end user experience

Objective 4: Engage and share risk intelligence with domestic and international regulatory partners and stakeholders

PRIORITIES

Objective 1:

- PPAD Adapt and leverage automation of licensing activities, reporting functions, directorate organizational charts and other administrative activities
- MDCP and CCBO Create and/or continue the implementation of the Transformation Roadmaps
- MDCP and CCBO Implement hybrid inspections where viable
- MDCP and CCBO Design an efficient and effective process for responding to inquiries for Border Operations and MDCP including general process improvements and the use of e-solutions (eg. Virtual assistant and/or automation of responses)
- CCBO Transform how BPCP delivers its core mandate by exploring suitability of conducting alternate compliance and enforcement activities to maintain oversight
- MDCP Explore the viability of issuing MD export certificates
- MDCP Enhance C&E approaches for non-compliant MDEL holders

Objective 2:

- PPAD Implement and update C&E tools (e.g. robust Site Risk Profiles)
- PPAD Ensure continued data analytics capacity
- PPAD and CCBO Explore the use of a "focused referral model" to address the constant backlog of referrals for Border Operations
- PPAD and MDCP Continue development of the risk intelligence tool, CONSILIUM, for Class II-IV manufacturers to strengthen post-market oversight

Objective 3:

- PPAD Contribute leadership and C&E perspective to the development of the Advanced Therapeutic Products (ATP) regulatory pathway
- PPAD and MDCP Support MDEL and Recall Amendments development and implementation (Phases 1 and 2)
- PPAD and CCBO Plan and implement the C&E components of Clinical Trial Modernization
- CCBO Improve BPCP's compliance promotion by proactively developing and sharing online key material for industry
- MDCP Scope and develop medical device shortages and exceptional importation program
- MDCP Develop and implement C&E strategy for mandatory incident reporting by hospitals
- MDCP Develop and implement C&E strategy for private label holders

Objective 1:

- PPAD Develop and implement an information management plan and strategy to securely store and access directorate data
- PPAD Develop and implement a centralised QMS function and dashboard by leading a data integration project to improve and streamline quality management
- PPAD Lead and promote quality management principles
- PPAD Support PRSD in the development and implementation of the ROEB FDA designation Directive and the certification standard
- PPAD Support the divisions in the implementation of the new Office 365 tools and Sharepoint

Objective 2:

- CCBO Determine the viability of centralizing the mailing of admissibility determination letters for Border Operations
- CCBO Develop program specific training modules for BPCP, CTCP and Border Operations
- MDCP Strengthen workload management of compliance verification incident reports
- MDCP Examine and strengthen risk-based approach to inspections

Objective 3:

- PPAD Support IT system development to replace the Inspection Reporting System (IRS) and eCES
- PPAD Develop interim IT solutions for Biological Products database (eg. IRS)
- PPAD Leverage the community practice to ensure access to new technology and IT (Power BI, Tableau, ROEB Data Lake, Tensor Flow, RPAs, etc.)
- CCBO Collaborate with HPCD and IMSD to update and further develop the Border Centres Operational Database (BCOD)
- MDCP Implement IT strategy for MDCP

Objective 4:

- PPAD and CCBO Enhance international partnerships with key regulators (eg. USFDA, MHRA, WHO)
- CCBO Increase transparency and information sharing of compliance and enforcement activities for CTCP with stakeholders and the public
- CCBO Proactively engage the CBSA operational senior management on emerging issues at the border and workload management
- CCBO Strengthen collaboration with HPFB partners
- MDCP Strengthen collaboration within Health Canada's medical devices partners (MDD)
- MDCP Regular information exchange with Medtech
- MDCP Establish regulatory engagement with US FDA Office of Regulatory Affairs

PILLARS

PEOPLE AND THE WORKPLACE

CREATE AND SUPPORT A RESPECTFUL, RESILIENT AND THRIVING WORKPLACE

OBJECTIVES

Objective 1: Continue to support employee's mental health and wellness

Objective 2: Foster an environment where employees are engaged and are provided with open, clear and transparent communication

Objective 3: Support a bilingual work environment where staff can work in the language of their choice.

Objective 4: Support employee career development and progress by providing fair, open and transparent opportunities and recognising and celebrating achievements

Objective 5: Provide timely access to mandatory training

PRIORITIES

Objective 1:

- DGO/Shared Priorities Implement the recommendations of the MDCCD Workplace Wellness Committee
- DGO/Shared Priorities Support the implementation of the ROEB Workplace Wellness Charter
- DGO/Shared Priorities Require employees to complete all corporate mandatory training (eg. workplace violence & harassment training)
- DGO/Shared Priorities Support employee re-entry in a workplace and ensure optimum communication between virtual and inperson staff

Objective 2:

- DGO Promote regular, meaningful interactions between senior management and employees
- DGO Issue the MDCCD Quarterly Newsletter to inform staff about developments, successes and employee movement
- MDCP and CCBO Build virtual team cohesion by supporting regular team meetings

Objective 3:

- DGO/Shared Priorities Ensure incumbents (managers and supervisors) meet positional language requirements
- DGO/Shared Priorities Integrate language training considerations into succession planning
- DGO/Shared Priorities Provide part-time language training opportunities for career development

Objective 4:

- DGO/Shared Priorities Focus on employee career development and retention
- DGO/Shared Priorities Maintain Succession Plan for key positions
- DGO/Shared Priorities Routinely recognize employee contributions, accomplishments and successes

Objective 5:

- DGO/Shared Priorities Support employee training and development
- PPAD Create an onboarding package for managers and employees