

RORB Directorates and corresponding Director Generals include:

### **Program Clusters**

- Health Product Compliance Directorate –Steve Schwendt, [Steven.Schwendt@hc-sc.gc.ca](mailto:Steven.Schwendt@hc-sc.gc.ca), (613)-957-6836
- Medical Devices and Clinical Compliance Directorate –Todd Cain, [Todd.Cain@hc-sc.gc.ca](mailto:Todd.Cain@hc-sc.gc.ca), (613)-668-2460

### **Delivery Clusters**

- Consumer Product Safety, Tobacco and Pesticide Directorate – Krista Locke, [Krista.Locke@hc-sc.gc.ca](mailto:Krista.Locke@hc-sc.gc.ca), (902)-407-7810
- Controlled Substances and Environmental Health Directorate – Ward Chickoski, [Ward.Chickoski@canada.ca](mailto:Ward.Chickoski@canada.ca), (780)-495- 3857
- Laboratories Directorate – Guy Aucoin, Guy [Aucoin@hc-sc.gc.ca](mailto:Aucoin@hc-sc.gc.ca) (450)-928-4100

### **Horizontal Clusters**

- Policy and Regulatory Strategies Directorate – Brenda Czich, [Brenda.Czich@hc-sc.gc.ca](mailto:Brenda.Czich@hc-sc.gc.ca), (613)-948-8431
- Planning and Operations Directorate – Scott McKenna, [Scott.McKenna@hc-sc.gc.ca](mailto:Scott.McKenna@hc-sc.gc.ca), (902)426-4600

Under this model, **Program Clusters** have been developed for health products (including drugs, over-the-counter drugs, biologics and radio-pharmaceuticals) and medical devices (including blood, donor semen, and cells, tissues and organs) where program design and compliance and enforcement activities will be managed in the same directorate. These clusters will continue to work closely with key functional areas in the Health Products and Food Branch.

The **Health Product Compliance Directorate** is responsible for the planning, delivery and issue resolution activities related to establishment licensing, Good Manufacturing Practices for health products (including drugs, over-the-counter drugs, biologics and radio-pharmaceuticals), drug compliance, verification and investigations and recalls, and Good Pharmacovigilance. It will also lead program direction and implementation for surveillance and risk intelligence, domestically and at the border.

The **Medical Devices and Clinical Compliance Directorate** is responsible for planning, delivery and issue resolution activities for medical device licensing and compliance verification. This unit will lead blood, tissue, organs, xenografts, good

## Appendix A

clinical practices and Canada Vigilance programs. It will also lead on recalls related to these product types and the border program for the branch.

**Delivery Clusters** have been developed for other Health Canada regulated products where only the focus will be primarily on compliance and enforcement, with strong alignment with the design element managed in partner branches, such as Healthy Environments and Consumer Product Safety (consumer products, controlled substances and tobacco), Pest Management Regulatory Agency (pesticides) and the Office of Medical Cannabis.

The **Horizontal Clusters** are key support areas that will provide policy and operational support for RORB.