A BRIEF NOTE ON REGULATORY REQUIREMENTS

The study of the Mid-Atlantic market for health care products does not include any reference to the regulatory requirements which must be met by exports to the U.S. This was done to focus the study on the commercial aspect of the market. A summary of these regulatory requirements may be found in a background paper entitled:

Summary of Regulatory Requirements for Medical Devices in Canada and the United States

Prepared by: Sector Analysis Division
Chemicals Branch
Department of Industry, Trade and Commerce
Ottawa, Canada

These papers are available from the Department of Industry, Trade and Commerce.