PrecisionBioLogic

Revised January 4, 2011

Subject: MSDS Information

Products marketed by Precision BioLogic Inc. are classified as Medical Devices within the Canadian Medical Devices Regulation and are, therefore, exempt from the Workplace Hazardous Material Information System (WHMIS) requirement to have a Material Safety Data Sheet provided.

Precision BioLogic products containing human blood have been manufactured with components that have been tested for the detection of hepatitis B surface antigen, antibodies to HIV and HCV, HIV-1 RNA, HCV RNA, and syphilis. All blood is collected in the US from FDA licensed centers and tested with FDA approved test kits. Additionally, the donors were screened for CJD/nvCJD and found acceptable.

Please note: No test method can provide total assurance that hepatitis B virus, human immunodeficiency virus, hepatitis C virus, or other infectious agents are absent.

Precision BioLogic supports safe laboratory practices and provides product handling instructions in the Direction Insert supplied with each product. Please refer to a product's Direction Insert for proper handling and disposal.

I trust that this information will fulfill your laboratory's safety requirements.

Sincerely,

Sandy Morrison
Director of Quality

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