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MATERIAL SAFETY DATA SHEET INFORMATION

Kerodex

All Medtech products, inclusive of the following brand name product, Kerodex, are registered with and defined by the United States Food and Drug Administration. Sec 1910.1200 of OSHA's Hazard Communication Standard states:

The purpose of this section is to ensure that the hazards of all chemicals produced or imported are evaluated and that information concerning their hazards is transmitted to employers and employees. This transmittal of information is to be accomplished by means of comprehensive hazard communication programs, which are to include container labeling and other forms of warning, material safety data sheets and employee training.

In reference to the above sections, the following:

Sec (5) This section does not require labeling, MSDS of any of the following chemicals: (ii) Any food, drug, cosmetic or medical device including materials intended for use as ingredients in such products as such terms are defined in the Federal Food and Drug and Cosmetic Act (21 U.S.C. 301 *et seq.*) and regulations issued under that, when they are subject to the labeling requirements under that Act by the Food and Drug Administration.

Sec (6) referring to Solid Waste Disposal Act amended by Resource Conservation and Recovery Act 1976 (42 U.S.C. 6901) (v) Food, drugs or cosmetics or alcoholic beverages in a retail establishment which are packaged for sale to consumers: and (vi) Food, drugs, cosmetics intended for personal consumption by employees in the work place; (viii) any drug, as that term is defined in the Federal Food, Drug and Cosmetic Act (21 U.S.C. 301 *et seq.*) when it is in solid, final form for direct administration to the patient (i.e. tablet or pills)."

And under SARA Title III: Emergency Planning and Community Right-To-Know Act 1986 (42 U.S.C. 11021) Sec 311 Material Safety Data sheets sec (e) "hazardous chemical" has the meaning given such term by section 1910.1200 (c) of Title 29 Code of Federal Regulations **except** that such term **does not include the following**:

- (1) Any food, food additive, color additive, **drug or cosmetic** regulated by the Food and Drug Administration;
- (2) Any substance to the extent it is used for personal, family or household purposes, or is present in the same form and concentration as a product packaged for distribution and use by the general public;
- (3) Any substance to the extent it is used in a research laboratory or other medical facility under the direct supervision of a technically qualified individual.

As defined by the above reference criterion, Kerodex 51 and Kerodex 71 do not require a Material Safety Data Sheet.