



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

AUG 14 2001

1375 '01 AUG 30 P2:08

Mr. Jack Speer  
Medical Instruments Technology, Inc.  
385 North 3050 East  
St. George, UT 84790

Re: P010010  
Reprocessed Cardiac Ablation Catheters

Dear Mr. Speer:

FDA received a citizen petition requesting an extension of enforcement discretion relating to the premarket approval (PMA) requirements under sections 513 and 515 of the Federal Food, Drug, and Cosmetic Act applicable to reprocessed single use devices. Previously, FDA issued a guidance on August 14, 2000 entitled "Enforcement Priorities for Single-Use Devices Reprocessed by Third Parties and Hospitals." This guidance stated that FDA intended to exercise enforcement discretion with respect to premarket submission requirements for reproprocessors of class III devices until August 14, 2001, provided that FDA received a PMA application no later than February 14, 2001, and FDA approval by August 14, 2001.

After considering the information in this petition, FDA believes that additional time is warranted, and is extending the timeframe for reproprocessors who have already filed PMAs for a period of 6 months to obtain FDA approval, and does not intend to enforce its premarket requirements with respect to those reproprocessors of single use class III devices until February 14, 2002. This extension, however, does not preclude the agency from taking immediate action if the agency becomes aware of additional risks posed by the product.

Sincerely Yours,

David W. Feigal, Jr., MD, MPH  
Director  
Center for Devices and Radiological Health

01P-0346

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