FDA 510(k) Premarket Notification

K071088 LIFE LATCH SHARPS & BIOHAZARD CONTAINER

CLEARED

Jul 5, 2007
78 days to decision

K071088 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k071088/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Apr 18, 2007
Decision date	Jul 5, 2007
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**M&M Industries, Inc.**
Location	Potomac, MD, US
Contact	NORMAN F ESTRIN
510(k) history	1 submissions · 1 cleared · 2007-2007

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k071088/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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