FDA 510(k) Premarket Notification

K951983 IMNI-II

CLEARED

Jun 20, 1995
54 days to decision

K951983 · Product code: **LRJ** · General Hospital
Source: https://www.510kdatabase.net/k951983/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Disinfectant, Medical Devices (LRJ)
Date received	Apr 27, 1995
Decision date	Jun 20, 1995
Days to decision	54 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cottrell, Ltd.**
Location	Englewood, CO, US
Contact	JACK SCOVILLE
510(k) history	9 submissions · 9 cleared · 1990-1999