FDA 510(k) Premarket Notification

K932922 PROCIDE 14 N.S., OMNICIDE/LIQUID & PLUS LIQ ACTIVATOR

CLEARED

Jul 3, 1995
748 days to decision

K932922 · Product code: **MED** · General Hospital
Source: https://www.510kdatabase.net/k932922/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilant, Medical Devices (MED)
Date received	Jun 15, 1993
Decision date	Jul 3, 1995
Days to decision	748 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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