

**K932922 PROCIDE 14 N.S., OMNICIDE/LIQUID & PLUS LIQ
ACTIVATOR**Jul 3, 1995
748 days to decisionK932922 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932922/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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