FDA 510(k) Premarket Notification

K933041 COLD CD ASP MODULE, ITE

CLEARED

Aug 2, 1993
96 days to decision

K933041 · Product code: **LRS** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k933041/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	I.v. Start Kit (LRS)
Date received	Apr 28, 1993
Decision date	Aug 2, 1993
Days to decision	96 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Electone, Inc.**
Location	Mchenry, IL, US
Contact	JIM NEE
510(k) history	73 submissions · 73 cleared · 1981-1997

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k933041/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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