FDA 510(k) Premarket Notification

K891023 SIEMENS CEILING STAND 800F, 800 AND 810

CLEARED

Jul 25, 1989
147 days to decision

K891023 · Product code: **LDQ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k891023/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Feb 28, 1989
Decision date	Jul 25, 1989
Days to decision	147 days
Third-party review	No

APPLICANT

Company	**Siemens Medical Electronics**
Location	Danvers, MA, US
Contact	MURFITT, PHD
510(k) history	15 submissions · 15 cleared · 1988-1995

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k891023/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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