FDA 510(k) Premarket Notification

K895676 HEMOSTAT/STIMULATOR PROBE

CLEARED

Dec 8, 1989
79 days to decision

K895676 · Product code: **ETN** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k895676/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve (ETN)
Date received	Sep 20, 1989
Decision date	Dec 8, 1989
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Rln Systems, Inc.**
Location	Jefferson City, MO, US
Contact	REA, M.D.
510(k) history	5 submissions · 5 cleared · 1989-2001

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k895676/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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