FDA 510(k) Premarket Notification

K871158 SIBIS REMOTE ACTUATOR

CLEARED

May 29, 1987
66 days to decision

K871158 · Product code: **HCB** · Neurology
Source: https://www.510kdatabase.net/k871158/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Aversive Conditioning (HCB)
Date received	Mar 24, 1987
Decision date	May 29, 1987
Days to decision	66 days
Third-party review	No

APPLICANT

Company	**Human Technologies, Inc.**
Location	St. Petersburg, FL, US
Contact	JAMES N DEERING
510(k) history	3 submissions · 3 cleared · 1987-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k871158/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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