FDA 510(k) Premarket Notification

K781623 PACING MODELS MUM 860, MUM 890

CLEARED

Dec 8, 1978
79 days to decision

K781623 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k781623/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Sep 20, 1978
Decision date	Dec 8, 1978
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**R.E. Brown Co., Inc.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1978-1978