FDA 510(k) Premarket Notification

K791380 VA 1000

CLEARED

Sep 19, 1979
57 days to decision

K791380 · Product code: **DTA** · Cardiovascular
Source: https://www.510kdatabase.net/k791380/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Pacemaker Electrode Function (DTA)
Date received	Jul 24, 1979
Decision date	Sep 19, 1979
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Vitatron Medical BV**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1976-1986