FDA 510(k) Premarket Notification

K760670 PACEMAKERS, VITATRONS (SERIES 2000)

CLEARED

Nov 12, 1976
56 days to decision

K760670 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k760670/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Sep 17, 1976
Decision date	Nov 12, 1976
Days to decision	56 days
Third-party review	No

APPLICANT

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