FDA 510(k) Premarket Notification

K900553 CONSTANT CURRENT STIMULATOR MODEL 022101

CLEARED

Mar 13, 1990
35 days to decision

K900553 · Product code: **JOQ** · Cardiovascular
Source: https://www.510kdatabase.net/k900553/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Pulse, Pacemaker, External Programmable (for Electrophysiological Studies Only) (JOQ)
Date received	Feb 6, 1990
Decision date	Mar 13, 1990
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Ep Technologies, Inc.**
Location	Mountain View, CA, US
Contact	S COOPERMAN
510(k) history	15 submissions · 15 cleared · 1988-2005