FDA 510(k) Premarket Notification

K905831 ARZCO OCTAPOLAR ESOPHAGEAL CARDIAC RECORD CATH

CLEARED

Mar 29, 1991
88 days to decision

K905831 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k905831/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Dec 31, 1990
Decision date	Mar 29, 1991
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Arzco Medical Electronics, Inc.**
Location	Vernon Hills, IL, US
Contact	RICHARD M BILOF
510(k) history	3 submissions · 3 cleared · 1991-1991