FDA 510(k) Premarket Notification

K900320 CRI CYNOSAR CATHETER

CLEARED

Sep 14, 1990
234 days to decision

K900320 · Product code: **DRA** · Cardiovascular
Source: https://www.510kdatabase.net/k900320/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Steerable (DRA)
Date received	Jan 23, 1990
Decision date	Sep 14, 1990
Days to decision	234 days
Third-party review	No

APPLICANT

Company	**Catheter Research C/O Burditt, Bowles & Radzius**
Location	Chicago, IL, US
Contact	BRIAN GRIGSBY
510(k) history	8 submissions · 8 cleared · 1987-1993