FDA 510(k) Premarket Notification

K894330 DAIG VESSEL DILATOR

CLEARED

Sep 12, 1989
60 days to decision

K894330 · Product code: **DRE** · Cardiovascular
Source: https://www.510kdatabase.net/k894330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Jul 14, 1989
Decision date	Sep 12, 1989
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**Daig Corp.**
Location	Mchenry, IL, US
Contact	J FLEISCHHACKER
510(k) history	63 submissions · 63 cleared · 1977-2000